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(54) Title: TETRAHYDROPTERIDINES AND PYRIDYLPYPERAZINES FOR TREATMENT OF NEUROLOGICAL DISORDERS		
(57) Abstract <p>Corticotropin releasing factor (CRF) antagonists of formula (I) and their use in treating anxiety, depression, and other psychiatric and neurological disorders.</p> <div data-bbox="1088 1113 1542 1323"><p style="text-align: right;">(I)</p></div>		

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TITLETetrahydropteridines and Pyridylpiperazines
for Treatment of Neurological Disorders

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FIELD OF THE INVENTION

This invention relates to compounds and pharmaceutical compositions, and to methods of using same in the treatment of psychiatric disorders and neurological diseases including major depression, anxiety-related disorders, post-traumatic stress disorder, supranuclear palsy and feeding disorders.

BACKGROUND OF THE INVENTION

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Corticotropin releasing factor (herein referred to as CRF), a 41 amino acid peptide, is the primary physiological regulator of proopiomelanocortin (POMC)-derived peptide secret. gland [J. Rivier et al., *Proc. Nat. Acad. Sci. (USA)* 80:4851 (1983); W. Vale et al., *Science* 213:1394 (1981)]. In addition to its endocrine role at the pituitary gland, immunohistochemical localization of CRF has demonstrated that the hormone has a broad extrahypothalamic distribution in the central nervous system and produces a wide spectrum of autonomic, electrophysiological and behavioral effects consistent with a neurotransmitter or neuromodulator role in brain [W. Vale et al., *Rec. Prog. Horm. Res.* 39:245 (1983); G.F. Koob, *Persp. Behav. Med.* 2:39 (1985); E.B. De Souza et al., *J. Neurosci.* 5:3189 (1985)]. There is also evidence that CRF plays a significant role in integrating the response of the immune system to physiological, psychological, and immunological stressors [J.E. Blalock, *Physiological Reviews* 69:1 (1989); J.E. Morley, *Life Sci.* 41:527 (1987)].

Clinical data provide evidence that CRF has a role in psychiatric disorders and neurological diseases including depression, anxiety-related disorders and feeding disorders. A role for CRF has also been postulated in the etiology and pathophysiology of Alzheimer's disease, Parkinson's disease, Huntington's disease, progressive supranuclear palsy and amyotrophic lateral sclerosis as they relate to the dysfunction of CRF neurons in the central nervous system [for review see E.B. De Souza, *Hosp. Practice* 23:59 (1988)].

In affective disorder, or major depression, the concentration of CRF is significantly increased in the cerebral spinal fluid (CSF) of drug-free individuals [C.B. Nemeroff et al., *Science* 226:1342 (1984); C.M. Banki et al., *Am. J. Psychiatry* 144:873 (1987); R.D. France et al., *Biol. Psychiatry* 28:86 (1988); M. Arato et al., *Biol Psychiatry* 25:355 (1989)]. Furthermore, the density of CRF receptors is significantly decreased in the frontal cortex of suicide victims, consistent with a hypersecretion of CRF [C.B. Nemeroff et al., *Arch. Gen. Psychiatry* 45:577 (1988)]. In addition, there is a blunted adrenocorticotropin (ACTH) response to CRF (i.v. administered) observed in depressed patients [P.W. Gold et al., *Am J. Psychiatry* 141:619 (1984); F. Holsboer et al., *Psychoneuroendocrinology* 9:147 (1984); P.W. Gold et al., *New Eng. J. Med.* 314:1129 (1986)]. Preclinical studies in rats and non-human primates provide additional support for the hypothesis that hypersecretion of CRF may be involved in the symptoms seen in human depression [R.M. Sapolsky, *Arch. Gen. Psychiatry* 46:1047 (1989)]. There is preliminary evidence that tricyclic antidepressants can alter CRF levels and thus modulate the numbers of CRF receptors in brain [Grigoriadis et al., *Neuropsychopharmacology* 2:53 (1989)].

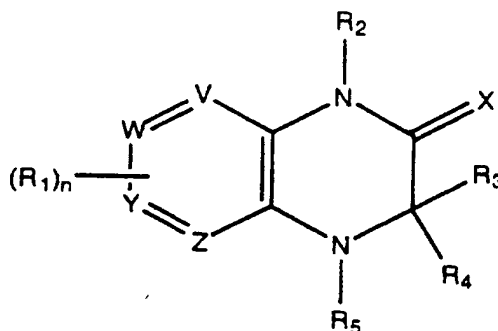
There has also been a role postulated for CRF in the etiology of anxiety-related disorders. CRF produces

anxiogenic effects in animals and interactions between benzodiazepine / non-benzodiazepine anxiolytics and CRF have been demonstrated in a variety of behavioral anxiety models [D.R. Britton et al., *Life Sci.* 31:363 (1982); C.W. Berridge and A.J. Dunn *Regul. Peptides* 16:83 (1986)]. Preliminary studies using the putative CRF receptor antagonist α -helical ovine CRF (9-41) in a variety of behavioral paradigms demonstrate that the antagonist produces "anxiolytic-like" effects that are qualitatively similar to the benzodiazepines [C.W. Berridge and A.J. Dunn *Horm. Behav.* 21:393 (1987), *Brain Research Reviews* 15:71 (1990)]. Neurochemical, endocrine and receptor binding studies have all demonstrated interactions between CRF and benzodiazepine anxiolytics providing further evidence for the involvement of CRF in these disorders. Chlordiazepoxide attenuates the "anxiogenic" effects of CRF in both the conflict test [K.T. Britton et al., *Psychopharmacology* 86:170 (1985); K.T. Britton et al., *Psychopharmacology* 94:306 (1988)] and in the acoustic startle test [N.R. Swerdlow et al., *Psychopharmacology* 88:147 (1986)] in rats. The benzodiazepine receptor antagonist (Ro15-1788), which was without behavioral activity alone in the operant conflict test, reversed the effects of CRF in a dose-dependent manner while the benzodiazepine inverse agonist (FG7142) enhanced the actions of CRF [K.T. Britton et al., *Psychopharmacology* 94:306 (1988)].

The mechanisms and sites of action through which the standard anxiolytics and antidepressants produce their therapeutic effects remain to be elucidated. It has been hypothesized however, that they are involved in the suppression of the CRF hypersecretion that is observed in these disorders. Of particular interest is that preliminary studies examining the effects of a CRF receptor antagonist (α -helical CRF9-41) in a variety of behavioral paradigms have demonstrated that the CRF

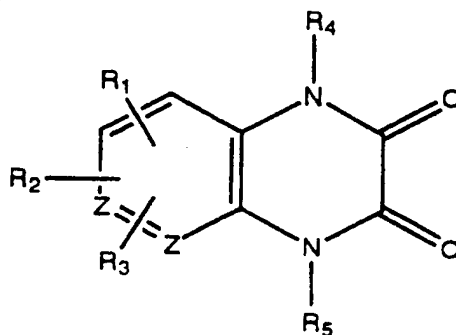
antagonist produces "anxiolytic-like" effects qualitatively similar to the benzodiazepines [for review see G.F. Koob and K.T. Britton, In: *Corticotropin-Releasing Factor: Basic and Clinical Studies of a Neuropeptide*, E.B. De Souza and C.B. Nemeroff eds., CRC Press p221 (1990)].

Several published patent applications disclose corticotropin releasing factor antagonist compounds. Among these are DuPont Merck PCT application US94/11050, Pfizer WO 95/33750, Pfizer WO 95/34563, and Pfizer WO 95/33727. U.S. Patent 5,424,311 discloses antiviral use of azaquinoxalines of the formula:



in which V, W, Y and Z are CH, CR¹, or N; X can be oxygen, sulfur or NR²; R¹ can be alkyl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, or alkylamino; R², R³, R⁴ and R⁵ can be hydrogen, alkyl, aryl or heteroaryl.

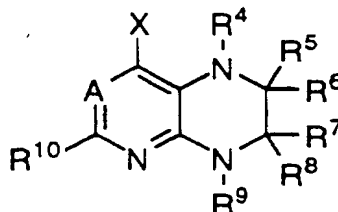
U.S. Patent 5,283,244 discloses glutamate receptor antagonizing activity of fused pyrazine derivatives of the the formula:



wherein Z represents C or N; R₁ represents a diazole or triazole substituent; and the other R groups represent hydrogen or various substituents such as alkyl, phenyl,
 5 or heterocycle.

SUMMARY OF THE INVENTION

This invention is a method of treating an affective
 10 disorder, anxiety, depression, irritable bowel syndrome, post-traumatic stress disorder, supranuclear palsy, immune suppression, Alzheimer's disease, gastrointestinal disease, anorexia nervosa or other feeding disorder, drug or alcohol withdrawal symptoms,
 15 drug addiction, or inflammatory disorder in a mammal comprising administering to the mammal a therapeutically effective amount of a CRF antagonist compound of formula I:



(I)

or a pharmaceutically acceptable salt or prodrug thereof, wherein:

- A is N or C-R¹¹;
 25 X is H, OR¹, S(O)_nR¹, NR¹R², CR¹R²R³, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈
 30 dialkylamino, or phenyl) or heteroaryl (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy,

- cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- n is 0, 1 or 2;
- R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, C₂-C₁₂ alkenyl, C₂-C₁₂ alkynyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, C₂-C₁₃ cyanoalkyl, C₂-C₅ carboalkoxy-(C₁-C₁₂ alkyl), phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl), or heteroaryl (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- R² and R³ are independently chosen from H, C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, C₂-C₁₂ alkenyl, C₂-C₁₂ alkynyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, C₂-C₁₃ cyanoalkyl, C₁-C₄ carboalkoxy, C₂-C₁₂ carboalkoxyalkyl, C(=O)CH₃, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl), or heteroaryl (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- R⁴ is H, C₁-C₁₂ alkyl, allyl, propargyl or benzyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl,

- nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- 5 R¹ and R⁴ may also optionally be taken together, along with the other four interconnected atoms, to form a ring of 5-9 total atoms, the structural sequence between the X group and the ring nitrogen atom consisting of the group (CH₂)_pW(CH₂)_q;
- p and q are independently 0, 1 or 2;
- 10 W is CH₂, C(CH₃)₂, C(=O), O, S or NCH₃;
- R⁵, R⁶, R⁷ and R⁸ are independently chosen from H, C₁-C₄ alkyl, allyl, propargyl, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl) or benzyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- 15 20 R⁴, R⁵ and R⁶ may also be taken together, along with the two interconnecting atoms, to constitute either an imidazole or tetrazole ring, the imidazole ring being optionally substituted with 1-2 groups chosen independently from C₁-C₄ alkyl or phenyl;
- 25 R⁵ and R⁶ may also be taken together to be O, S or NR¹²;
- 30 R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), pyridyl (optionally substituted with 1-4 groups chosen from halogen,
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- C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), or pyrimidyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano);
- R¹⁰ is H, C₁-C₄ alkyl or cyano;
- R¹¹ is H, C₁-C₄ alkyl or halogen;
- R¹² is H, C₁-C₄ alkyl or phenyl;
- aryl is phenyl, biphenyl or naphthyl; and heteroaryl is pyridyl, pyrimidinyl, triazinyl, furanyl, quinolinyl, isoquinolinyl, thienyl, imidazolyl, thiazolyl, indolyl, pyrrolyl, oxazolyl, benzofuranyl, benzothienyl, benzthiazolyl, isoxazolyl or pyrazolyl.

- Compounds of formula I, other than those in which R₅ and R₆ are taken together and are O, S or NR₁₂, are novel. This invention includes the novel compounds of formula I and pharmaceutical compositions containing them.

- Preferred compounds for use in the method of this invention are compounds of formula (I) wherein:

- X is OR¹, NR¹R², CR¹R²R³, or phenyl (optionally substituted at the 2-position with CF₃, nitro, halogen or cyano);
- R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, or phenyl (optionally substituted with 1-4 groups independently chosen from halogen, haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH,

- C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
- R⁴ is H or C₁-C₄ alkyl;
- R⁵ and R⁶ are either H or C₁-C₄ alkyl;
- 5 R⁴, R⁵ and R⁶ may also be taken together, along with the two interconnecting atoms, to constitute a tetrazole ring;
- R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), 3-pyridyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), or 5-pyrimidyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano);
- 10 20 R¹⁰ is CH₃;
- and R¹¹ is H.
- 25 More preferred compounds in this invention are of the formula (I) wherein:
- A is N;
- X is NR¹R² or CR¹R²R³;
- R¹ is C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;
- 30 R² and R³ are independently H, C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;
- R⁴ is H;
- R⁵ and R⁶ are H;
- R⁷ and R⁸ are independently H or CH₃;
- 35 and R⁹ is phenyl (optionally substituted with 1-4 groups

chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano).

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Specifically preferred because of their biological activity are the following compounds:

- 8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;
- 10 8-(2-chloro-4,6-dimethoxyphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;
- 4-(ethylbutylamino)-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine;
- and 4-(1-methoxy-2-butyl)amino-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine.
- 15

DETAILED DESCRIPTION OF THE INVENTION

- Many compounds of this invention have one or more asymmetric centers or planes. Unless otherwise indicated, all chiral (enantiomeric and diastereomeric) and racemic forms are included in the present invention. Many geometric isomers of olefins, C=N double bonds, and the like can also be present in the compounds, and all such stable isomers are contemplated in the present invention. The compounds may be isolated in optically active or racemic forms. It is well known in the art how to prepare optically active forms, such as by resolution of racemic forms or by synthesis from optically active starting materials. All chiral, (enantiomeric and diastereomeric) and racemic forms and all geometric isomeric forms of a structure are intended, unless the specific stereochemistry or isomer form is specifically indicated.
- 35 The term "alkyl" includes both branched and straight-chain alkyl having the specified number of carbon atoms. "Alkenyl" includes hydrocarbon chains of

either a straight or branched configuration and one or more unsaturated carbon-carbon bonds which may occur in any stable point along the chain, such as ethenyl, propenyl, and the like. "Alkynyl" includes hydrocarbon chains of either a straight or branched configuration and one or more triple carbon-carbon bonds which may occur in any stable point along the chain, such as ethynyl, propynyl and the like. "Haloalkyl" is intended to include both branched and straight-chain alkyl having the specified number of carbon atoms, substituted with 1 or more halogen; "alkoxy" represents an alkyl group of indicated number of carbon atoms attached through an oxygen bridge; "cycloalkyl" is intended to include saturated ring groups, including mono-, bi- or poly-cyclic ring systems, such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and so forth. "Halo" or "halogen" includes fluoro, chloro, bromo, and iodo.

The term "substituted", as used herein, means that one or more hydrogen on the designated atom is replaced with a selection from the indicated group, provided that the designated atom's normal valency is not exceeded, and that the substitution results in a stable compound. When a substituent is keto (i.e., =O), then 2 hydrogens on the atom are replaced.

Combinations of substituents and/or variables are permissible only if such combinations result in stable compounds. By "stable compound" or "stable structure" is meant a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into an efficacious therapeutic agent.

The term "appropriate amino acid protecting group" means any group known in the art of organic synthesis for the protection of amine or carboxylic acid groups. Such amine protecting groups include those listed in Greene and Wuts, "Protective Groups in Organic Synthesis" John Wiley & Sons, New York (1991) and "The

Peptides: Analysis, Synthesis, Biology, Vol. 3, Academic Press, New York (1981), the disclosure of which is hereby incorporated by reference. Any amine protecting group known in the art can be used. Examples of amine protecting groups include, but are not limited to, the following: 1) acyl types such as formyl, trifluoroacetyl, phthalyl, and p-toluenesulfonyl; 2) aromatic carbamate types such as benzyloxycarbonyl (Cbz) and substituted benzyloxycarbonyls, 1-(p-biphenyl)-1-methylethoxycarbonyl, and 9-fluorenylmethyloxycarbonyl (Fmoc); 3) aliphatic carbamate types such as tert-butyloxycarbonyl (Boc), ethoxycarbonyl, diisopropylmethoxycarbonyl, and allyloxycarbonyl; 4) cyclic alkyl carbamate types such as cyclopentyloxycarbonyl and adamantyloxycarbonyl; 5) alkyl types such as triphenylmethyl and benzyl; 6) trialkylsilane such as trimethylsilane; and 7) thiol containing types such as phenylthiocarbonyl and dithiasuccinoyl.

The term "amino acid" as used herein means an organic compound containing both a basic amino group and an acidic carboxyl group. Included within this term are natural amino acids, modified and unusual amino acids, as well as amino acids which are known to occur biologically in free or combined form but usually do not occur in proteins. Included within this term are modified and unusual amino acids, such as those disclosed in, for example, Roberts and Vellaccio (1983) The Peptides, 5: 342-429, the teaching of which is hereby incorporated by reference. Modified or unusual amino acids which can be used to practice the invention include, but are not limited to, D-amino acids, hydroxylysine, 4-hydroxyproline, an N-Cbz-protected amino acid, ornithine, 2,4-diaminobutyric acid, homoarginine, norleucine, N-methylaminobutyric acid, naphthylalanine, phenylglycine, β -phenylproline, tert-leucine, 4-aminocyclohexylalanine, N-methyl-

norleucine, 3,4-dehydroproline, N,N-dimethylaminoglycine, N-methylaminoglycine, 4-aminopiperidine-4-carboxylic acid, 6-aminocaproic acid, trans-4-(aminomethyl)-cyclohexanecarboxylic acid, 2-, 3-, and 4-(aminomethyl)-benzoic acid, 1-aminocyclopentanecarboxylic acid, 1-aminocyclopropanecarboxylic acid, and 2-benzyl-5-aminopentanoic acid.

The term "amino acid residue" as used herein means that portion of an amino acid (as defined herein) that is present in a peptide.

The term "peptide" as used herein means a compound that consists of two or more amino acids (as defined herein) that are linked by means of a peptide bond. The term "peptide" also includes compounds containing both peptide and non-peptide components, such as pseudopeptide or peptide mimetic residues or other non-amino acid components. Such a compound containing both peptide and non-peptide components may also be referred to as a "peptide analog".

The term "peptide bond" means a covalent amide linkage formed by loss of a molecule of water between the carboxyl group of one amino acid and the amino group of a second amino acid.

The term "pharmaceutically acceptable salts" includes acid or base salts of the compounds of formulas (I) and (II). Examples of pharmaceutically acceptable salts include, but are not limited to, mineral or organic acid salts of basic residues such as amines; alkali or organic salts of acidic residues such as carboxylic acids; and the like.

Pharmaceutically acceptable salts of the compounds of the invention can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl

acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in Remington's Pharmaceutical Sciences, 17th ed., Mack Publishing Company, Easton, PA, 1985, p. 1418, the disclosure of which is hereby incorporated by reference.

"Prodrugs" are considered to be any covalently bonded carriers which release the active parent drug of formula (I) or (II) *in vivo* when such prodrug is administered to a mammalian subject. Prodrugs of the compounds of formula (I) and (II) are prepared by modifying functional groups present in the compounds in such a way that the modifications are cleaved, either in routine manipulation or *in vivo*, to the parent compounds. Prodrugs include compounds wherein hydroxy, amine, or sulfhydryl groups are bonded to any group that, when administered to a mammalian subject, cleaves to form a free hydroxyl, amino, or sulfhydryl group, respectively. Examples of prodrugs include, but are not limited to, acetate, formate and benzoate derivatives of alcohol and amine functional groups in the compounds of formulas (I) and (II); and the like.

The term "therapeutically effective amount" of a compound of this invention means an amount effective to antagonize abnormal level of CRF or treat the symptoms of affective disorder, anxiety or depression in a host.

Synthesis

Synthesis of compounds of Formula (I) wherein A = N may begin with amidine compounds of Formula (II) (Scheme I), which are available commercially or synthetically from heating a nitrile compound and an ammonium salt. Compound (II) may then be condensed with a malonate ester (using conditions such as sodium in ethanol) to give a dihydroxy-pyrimidine compound of Formula (III). Nitration at the 5-position may be accomplished through the use of such conditions as concentrated nitric acid

with or without the presence of another acid such as concentrated sulfuric or glacial acetic. The hydroxy groups of the nitrated compound of Formula (IV) may then be converted into leaving groups (Y), which include

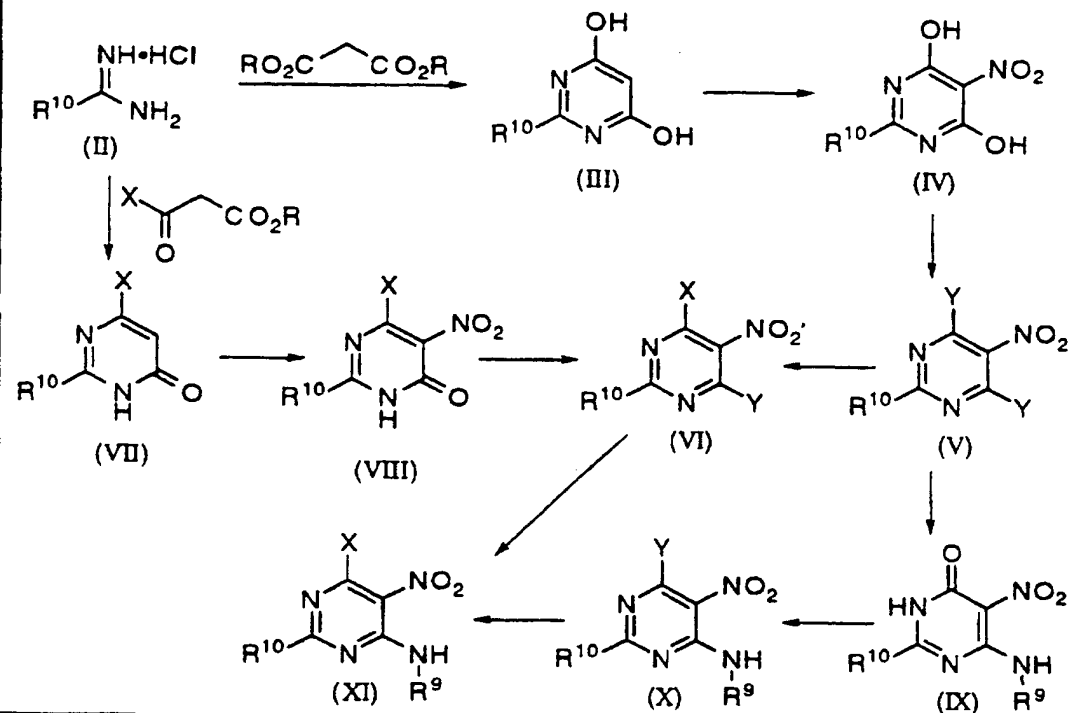
5 chloro, bromo, toluenesulfonate, or methanesulfonate. The dichloro compound (Formula (V), $Y = Cl$) may be prepared from the dihydroxy by a reagent such as phosphorus oxychloride, with or without the assistance of a catalyst such as diethylaniline. The

10 bis(toluenesulfonate) compound (Formula (V), $Y = OSO_2C_6H_4CH_3$), may be prepared from the dihydroxy compound by treatment with a reagent such as toluenesulfonic anhydride. Careful addition one

15 equivalent of a suitable form of a compound X-H to the compound of Formula (V) results in replacement of one of the Y groups with X. This is of particular utility when the X group represents a nucleophilic atom, such as nitrogen, sulfur or oxygen. Conditions which will facilitate this transformation include the optional

20 presence of bases such as sodium hydride, triethylamine, diisopropylethylamine or

Scheme I



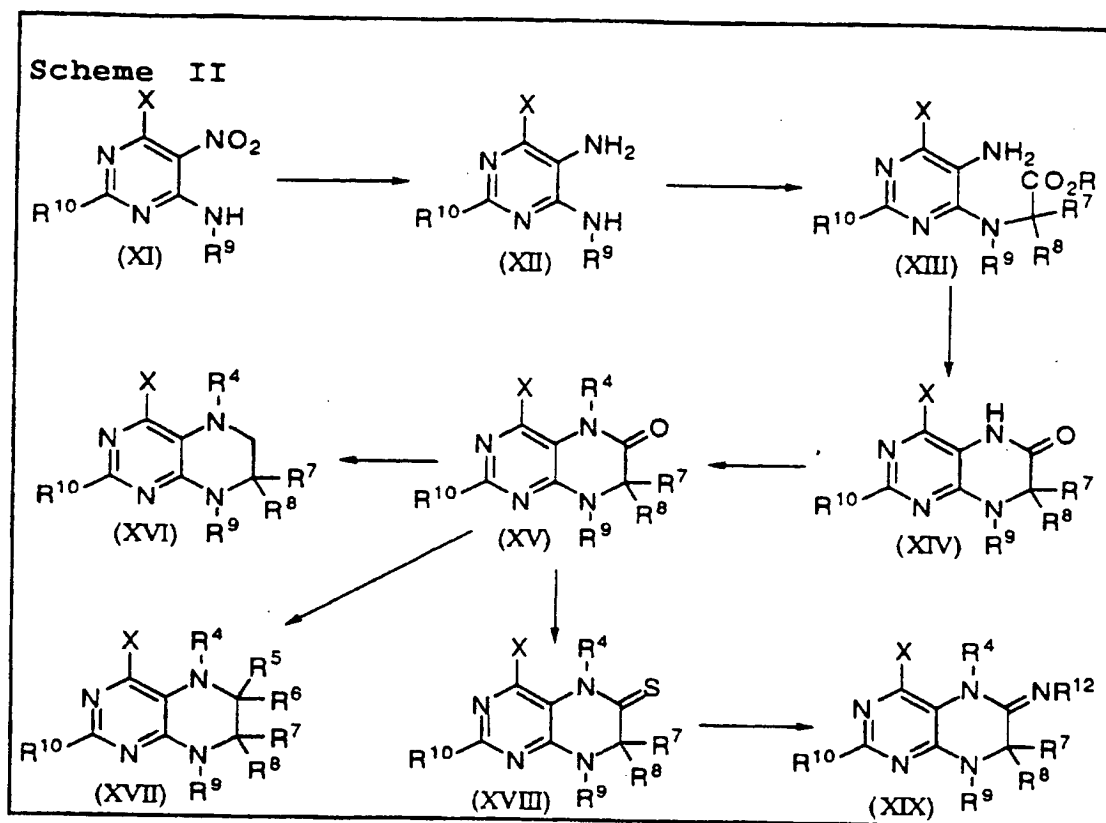
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potassium carbonate, in solvents such as tetrahydrofuran, dimethylformamide, dimethylsulfoxide, methylene chloride, acetonitrile or ethanol, at appropriate temperatures.

Alternatively, in the case where X represents a group without a corresponding nucleophilic compound X-H being available, one may condense a compound of Formula (II) with an appropriately-substituted ketoester (using conditions similar to those for the malonate condensation) to obtain a compound of Formula (VII). Nitration conditions similar to those described above may then be used to prepare the nitro compound (VIII). Conversion of the pyrimidone group to the desired Y group may then be accomplished using the same conditions as described above for the transformation of (IV) to (V).

20

- A third alternative involves treatment of the compound of Formula (V) with a compound R^9-NH_2 . Conditions may be found for each Y group so that one Y group is replaced by R^9-NH , and the other is hydrolyzed to the pyrimidone (compound Formula (IX)). For example, for $Y = Cl$, this conversion may be effected by slow addition of a dimethylsulfoxide solution of one equivalent of R^9-NH_2 to a dimethylsulfoxide solution of compound (V), followed by aqueous workup. The pyrimidone of Formula (IX) may be converted to Y-bearing compound (Formula (X)) using the conditions described above for (IV) to (V). The Y group can then be replaced with X analogously to the transformation of (V) to (VI) to give a compound of Formula (XI).
- Alternatively, the compound of Formula (VI) may be converted to the compound of Formula (XI) by treatment with the compound R^9-NH_2 . Suitable conditions for this reaction include treatment with excess sodium hydride in refluxing toluene or heating the two compounds together in an alcoholic solvent (ethanol, propanol, butanol, ethylene glycol, ethoxyethoxyethanol) or other polar, aprotic solvents (such as dimethylformamide, 1,4-dioxane, dimethoxyethane or diglyme) without a base to effect the coupling.
- Scheme II shows the appending of the second ring onto the pyrimidine ring. The nitro group in the compound of Formula (XI) can be reduced to an amino group using conditions such as sodium dithionite, catalytic hydrogenation, iron or zinc. The compound of Formula (XII) may be treated with a base such as sodium hydride (in solvents such as dimethylformamide, dimethyl sulfoxide, 1,4-dioxane, tetrahydrofuran, etc.), followed by a reagent of the general formula $Y-CR^7R^8-CO_2R$, where Y is halogen or pseudohalogen, and the structure of R is only important if



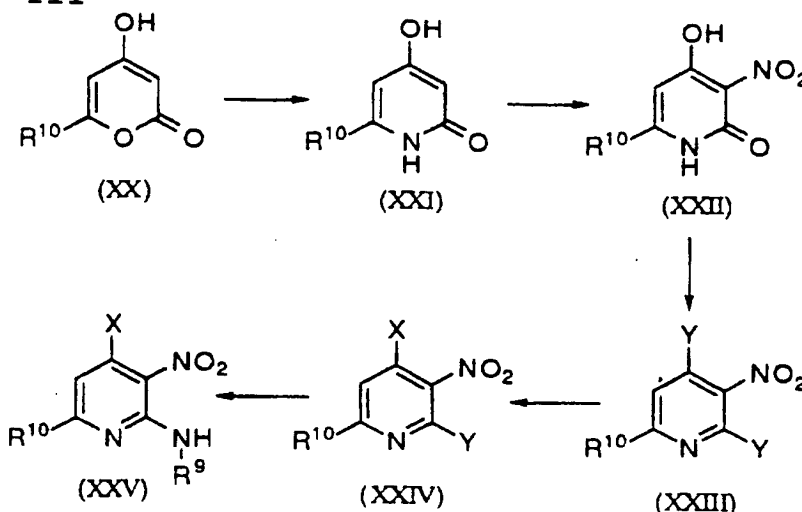
- 5 removal of the group prior to cyclization is desired. Cyclization of the compound of Formula (XIV) may be accomplished by heating in a solvent such as ethanol, dimethylformamide, etc. at temperatures ranging anywhere from ambient to the boiling point of the solvent. An
- 10 additive such as an acid source (such as toluenesulfonic acid, aqueous hydrochloric, etc.), a base (triethylamine, sodium hydroxide, etc.) or a physical catalyst (such as molecular sieves) may be added, in quantities ranging from catalytic to stoichiometric to
- 15 excess. In practice, the cyclization of (XIII) often is very facile, particularly in the case where R is lower alkyl, and will occur spontaneously in the reaction medium of the alkylation of compound (XII). Cyclized compound (XIV) may be alkylated with the R⁴ group by
- 20 first treatment with a base such as sodium hydride in a solvent such as dimethylformamide or dimethylsulfoxide, then an alkylating reagent (such as a halogen- or pseudohalogen-bearing compound) which provides the R⁴

group, to provide the compound of Formula (XV). At this point, compounds derived from bromoacetate alkylation of compound (XII) can be alkylated with appropriate R^7 and R^8 by treatment with a strong base such as sodium
5 hydride, lithium diisopropylamide or sodium hexamethyldisilazide, and then alkylating agents bearing the R^7 or R^8 groups, thus resulting in the compound of Formula (XV).

Compound (XV) is a key intermediate which may be
10 used to generate variations of Formula (I). For example, the carbonyl group of compound (XV) may be reduced with reagents such as lithium aluminum hydride, borane (complexed with tetrahydrofuran or other suitable ligands) or diisobutylaluminum hydride, which will
15 generate a compound of Formula (XVI). The carbonyl group may be substituted with R^5 and R^6 groups using appropriately-substituted organolithium or organomagnesium reagents, to prepare compounds of Formula (XVII). The carbonyl group of compound (XV) may
20 be converted to thiocarbonyl by treatment with reagents such as Lawesson's Reagent or phosphorus pentasulfide in appropriate solvents (toluene, benzene, etc.). The thioamide group of compound (XVIII) may be converted to amidine using the method of Robba et al. (*Tetrahedron*
25 *Letters* 1992, 33, 2803-2804), which involves treatment with an amine of formula $R^{11}-NH_2$ and a catalyst such as a mercury (II) salt. This will result in the synthesis of a compound of Formula (XIX).

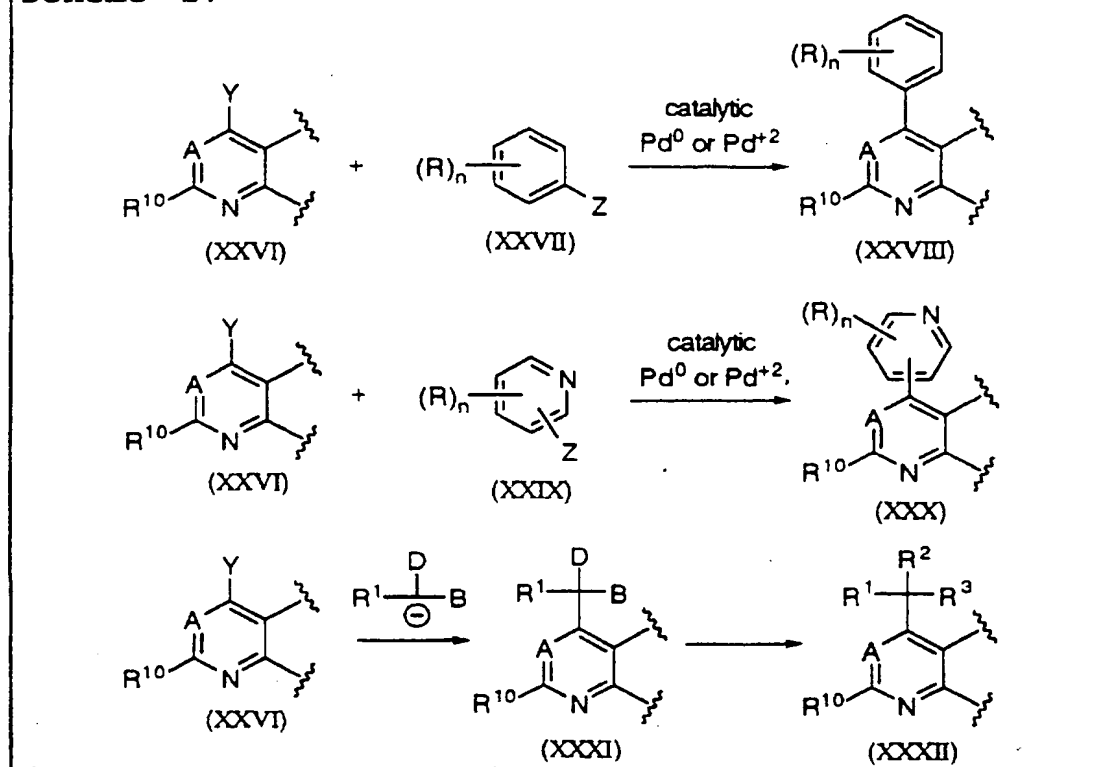
Compounds of Formula (I) composed of a fused
30 pyridine ring ($A = CH$) may be prepared using very similar technology to that presented in Scheme II. In this case, however, the starting material is not of the structure (XI), but rather

Scheme III



- 5 of structural formula (XXV) (Scheme III). This compound may be prepared starting with a lactone compound of Formula (XX), which are available by dimerization of a ketoester $R^{10}C(=O)CH_2CO_2Et$ according to the method of Arndt (*Org. Syn., Coll. Vol. III*, p. 231), followed by
- 10 deacylation according to the method of Collie et al. (*J. Chem. Soc.* **1907**, 91, p. 787 and references therein). The ring oxygen atom may be replaced with nitrogen by treatment with conc. aq. ammonium hydroxide, according to the method of Wang (*J. Heterocyclic Chem.* **1970**, 1, 389-392).
- 15 Compound (XXI) may be nitrated similarly to the transformation of compound (III) to give compound (XXII). The hydroxy groups of compound (XXII) may be converted to leaving groups Y using the techniques discussed above for the conversion of compound (IV) to (V).
- 20 The C⁴ Y group may be selectively replaced with a nucleophilic X group, and the other Y group in compound (XXIV) may be replaced with NHR^9 by treatment with a compound R^9NH_2 , either with no solvent or an appropriate solvent (such as a high-boiling alcohol) at temperatures sufficiently elevated to effect coupling. Compound (XXV)
- 25

Scheme IV



5 may then be employed in the same general way as for compound (XI) to generate compounds of Formula (I).

Further functionalization of this class of compounds may be achieved using a compound of Formula (XXVI) (Scheme IV), which represents some pyridine or pyridine compound (either uncyclized, like compounds (XI) or (XXV), or a cyclized compound) bearing a leaving group Y. The Y group may be replaced with phenyl or pyridyl using coupling reactions employing a phenyl (or pyridyl) compound of Formula (XXVII) (or (XXIX)) and an appropriate palladium catalyst. For example, arylboronic acids ($\text{Z} = \text{B}(\text{OH})_2$) may be coupled to a heterocyclic halide using catalytic amounts of tetrakis(triphenylphosphine)palladium, which is the method of Suzuki, et al. (*Synthetic Communications* 1981, 11, p. 513-519). Other appropriate reagents for this coupling reaction includes organomagnesium ($\text{Z} = \text{MgBr}$ or MgCl) reagents (with nickel (II) chloride

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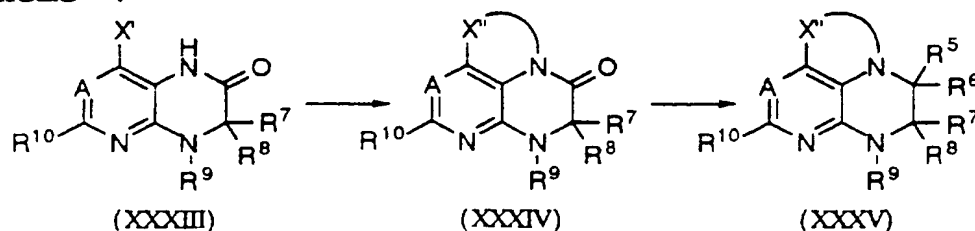
catalysis according to the method of Sugimori et al.,
Synthetic Communications **1991**, 21, p. 481-487) or
organozinc (Z = ZnCl) reagents (according to the method
of Negishi et al., *J. Org. Chem.* **1977**, 42, p. 1821-
5 1823).

Other carbon substituents may be introduced into
compound (XXVI) by treatment with a sodium salt
(generated by the use of a base such as sodium ethoxide
or sodium hydride) of an active methylene or methine
10 reagent (i.e. where B and D are groups which stabilize
adjacent anions, such as keto, carboalkoxy, cyano,
alkyl- or aryl-sulfonyl, etc.). The resulting compounds
of Formula (XXXI) may be further modified by conversion
of the B and D groups into R² and R³ groups. Those
15 skilled in the art of organic synthesis should readily
understand possible variations of these conversions to
prepare a number of different R¹, R² and R³ group
substituents.

Preparation of compounds of Formula (I) wherein the
20 R¹ and R⁴ groups are taken together to form a ring may
be accomplished beginning from a compound of Formula
(XXXIII) (Scheme V), where X' is meant to designate a
group NHR², OH, SH or CHR²R³. This compound may be
treated with a base (such as sodium hydride) in an
25 appropriate solvent, followed by a reagent bearing
reactive terminii on both ends (for example, a
dihaloalkane, a haloester, etc.). The X' and amide NH
groups will couple with such a reagent under these
conditions to form the third ring of compound (XXXIV).
30 The amide group may then be modified as described above
to give then final product of Formula (XXXV).

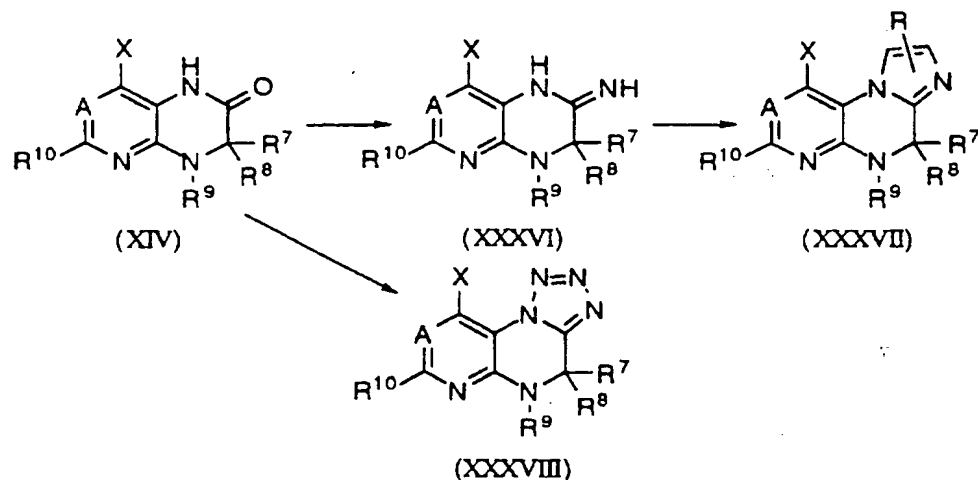
Compounds of Formula (I) wherein the R⁴, R⁵ and R⁶
groups are taken together to form a heteroaromatic ring
may

Scheme V



- 5 be prepared using the strategy displayed in Scheme VI. Compound (XIV) may be converted to amidine (XXXVI), using the conditions described above for the preparation of compound (XIX). The amidine is treated with an α -halo- or α -hydroxyketone, under conditions such as
- 10 refluxing alcohol, to afford the imidazole compound (XXXVII). Compound (XIV) may be converted to fused tetrazole compound (XXXVIII) using the conditions of Duncia et al. (*J. Org. Chem.* **1991**, 56, p. 2395).

15 Scheme VI



The experimental methods listed below for Examples 1, 17, 24, 42, 131, 143, 155, and 248 may be used in the preparation of all the compounds shown in Tables I (pyrimidines) and II (pyridines).

5

Example 1

Preparation of 8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridin-6-one

- 10 **Part A.** A solution of 4,6-dichloro-2-methyl-5-nitropyrimidine (prepared using the methods of Albert, et al., *J. Chem. Soc.* **1954**, p. 3832) (2.77 g, 13.3 mmol) in absolute ethanol (25 mL) was cooled to 0 °C, and treated with triethylamine (2.00 mL, 14.3 mmol).
- 15 Then, a solution of ethylbutylamine (1.80 mL, 13.2 mmol) in ethanol (3 mL) was added dropwise with stirring. The mixture was allowed to stir and warm to ambient temperature overnight, then was partitioned between water and ethyl acetate (100 mL each). The organic
- 20 phase was separated, washed with satd. aq. brine (100 mL), dried over anhydrous sodium sulfate, filtered and evaporated. The residue was separated by column chromatography (silica gel, dichloromethane) to afford 4-chloro-6-(ethylbutylamino)-2-methyl-5-nitropyrimidine
- 25 as an oil (3.34 g, 12.2 mmol, 92%). Spectral data: TLC R_f 0.59 (dichloromethane). ^1H NMR (300 MHz, CDCl_3): δ 3.45 (2H, q, $J = 7.0$ Hz), 3.38 (2H, t, $J = 7.7$ Hz), 2.50 (3H, s), 1.62-1.52 (2H, m), 1.38-1.26 (2H, m), 1.20 (3H, t, $J = 7.0$ Hz), 0.94 (3H, t, $J = 7.3$ Hz). MS ($\text{NH}_3\text{-CI}$):
- 30 m/e 276 (4), 275 (40), 274 (16), 273 (100).

- Part B.** The product of Part A above (2.97 g, 10.9 mmol) was dissolved in ethoxyethoxyethanol solution (11 mL), and treated with 2-bromo-4-isopropylaniline (2.34 g, 10.9 mmol). The mixture was heated to 120 °C for 4
- 35 h, then cooled and partitioned between water and ethyl acetate (100 mL each). The organic layer was separated and washed with two additional portions of water (100 mL

each) and brine (100 mL). The aqueous phases were all back-extracted in sequence with more ethyl acetate (100 mL). The extracts were combined, dried over sodium sulfate, filtered and evaporated. The residue was separated by column chromatography (silica gel, 5:95 ethyl acetate-hexane) to afford 6-(2-bromo-4-isopropylphenylamino)-4-(ethylbutylamino)-2-methyl-5-nitropyrimidine as an oil (3.05 g, 6.77 mmol, 62%). Spectral data: TLC R_f 0.56 (20:80 ethyl acetate-hexane). ^1H NMR (300 MHz, CDCl_3): δ 10.30 (1H, br s), 8.32 (1H, d, $J = 8.4$ Hz), 7.45 (1H, d, $J = 1.8$ Hz), 7.19 (1H, dd, $J = 8.4, 1.8$ Hz), 3.52-3.42 (4H, m), 2.88 (1H, heptet, $J = 7.0$ Hz), 2.40 (3H, s), 1.70-1.58 (2H, m), 1.39-1.29 (2H, m), 1.26 (3H, t, obscured), 1.25 (6H, d, $J = 7.0$ Hz), 0.94 (3H, t, $J = 7.1$ Hz). MS ($\text{NH}_3\text{-CI}$): m/e 454 (4), 453 (24), 452 (100), 451 (26), 450 (99).

Part C. The product of Part B above (256 mg, 568 μmol) was dissolved in 1:1 dioxane-water (3 mL), and treated with conc. aq. ammonium hydroxide (0.5 mL). To this was added, with stirring, solid sodium dithionite (440 mg, 2.53 μmol) in 3 portions over 1 h. The resulting solution was allowed to stir for an additional 8 h, then partitioned between water and ethyl acetate (100 mL). The organic layer was washed with brine, dried over sodium sulfate, filtered and evaporated. The residual material was purified by elution through a short plug of silica gel (20:80 ethyl acetate-hexane), and evaporation gave pure 5-amino-6-(2-bromo-4-isopropylphenylamino)-4-(ethylbutylamino)-2-methylpyrimidine as an oil (198 mg, 472 μmol , 83%). Spectral data: TLC R_f 0.26 (10:90 ethyl acetate-hexane). ^1H NMR (300 MHz, CDCl_3): δ 8.15 (1H, d, $J = 8.4$ Hz), 7.38 (1H, d, $J = 2.2$ Hz), 7.15 (1H, dd, $J = 8.4, 2.2$ Hz), 7.04 (1H, br s), 3.30-3.19 (4H, m), 3.08 (2H, br s), 2.85 (1H, heptet, $J = 7.0$ Hz), 2.47 (3H, s), 1.57-1.44 (2H, m), 1.39-1.26 (2H, m), 1.23 (6H, d, $J = 7.0$ Hz), 1.11 (3H, t, $J = 7.0$ Hz), 0.91 (3H, t, $J = 7.1$ Hz). MS

(NH₃-CI): m/e 424 (3), 423 (24), 422 (100), 421 (26), 420 (100).

Part D. Sodium hydride dispersion in mineral oil (0.26 g w/w, 5.42 mmol) was washed with hexane, and the hexane was decanted off. The remaining solid was dried under vacuum and suspended in anhydrous dimethylformamide (5 mL). The resulting suspension was cooled in an ice bath, while the product of Part C above (1.78 g, 4.23 mmol) in dimethylformamide solution (5 mL) was added slowly by syringe. After hydrogen evolution was complete, the mixture was treated slowly with ethyl bromoacetate (0.47 mL, 4.24 mmol) by syringe, and the mixture was allowed to stir for 10 h. It was partitioned between water and ethyl acetate (100 mL each), and the organic phase was washed with two additional portions of water (100 mL each) and brine (100 mL). The aqueous phases were back-extracted in sequence with ethyl acetate (100 mL), and the extracts were combined, dried over sodium sulfate, filtered and evaporated. The residue was separated by column chromatography (silica gel, 10:90 ethyl acetate-hexane) to afford the title product as a solid (1.35 g, 2.93 mmol, 69%). Spectral data: m.p. 146-147 °C. TLC R_f 0.49 (50:50 ethyl acetate-hexane). ¹H NMR (300 MHz, CDCl₃): δ 7.53 (1H, s), 7.36 (1H, br s), 7.25 (2H, s), 4.44 (1H, d, J = 15 Hz), 4.25 (1H, d, J = 15 Hz), 3.29-3.15 (4H, m), 2.94 (1H, heptet, J = 7.0 Hz), 2.29 (3H, s), 1.54-1.45 (2H, m), 1.39-1.29 (2H, m), 1.28 (6H, d, J = 7.0 Hz), 1.11 (3H, t, J = 7.1 Hz), 0.92 (3H, t, J = 7.1 Hz). MS (NH₃-CI): m/e 464 (3), 463 (24), 462 (100), 461 (26), 460 (99). Analysis calc'd for C₂₂H₃₀BrN₅O: C, 57.39; H, 6.58; N, 15.21; found: C, 56.74; H, 6.02; N, 14.41.

Examples 143 and 155

Preparation of 8-(2-bromo-4-isopropylphenyl)-2,5-dimethyl-4-(ethylbutylamino)-5,6,7,8-tetrahydropteridin-6-one and 8-(2-bromo-4-isopropylphenyl)-4-

(ethylbutylamino)-2,5,7-trimethyl-5,6,7,8-tetrahydropteridin-6-one

Part A. Sodium hydride suspension in mineral oil (200 mg of 50% w/w, 4.17 mmol) was washed with hexane, dried under vacuum and suspended in anhydrous dimethylformamide (5 mL). This suspension was cooled to 0 °C, while a solution of the compound of Example 1 (1.58 g, 3.43 mmol) in dimethylformamide (5 mL) was added slowly by syringe. The resulting mixture was allowed to stir for 1 h, then was treated with methyl iodide (0.30 mL, 4.82 mmol) by syringe. The mixture was allowed to stir overnight, then was partitioned between water and ethyl acetate (100 mL each). The organic phase was washed with two additional portions of water and one of brine solution. The aqueous fractions were back-extracted in sequence with more ethyl acetate, and the organic layers were combined, dried over sodium sulfate, filtered and evaporated. The residual oil was separated by column chromatography (silica gel, 10:90 ethyl acetate-hexane) to afford two fractions. The first compound off the column was the title compound of Example 155 (150 mg, 0.31 mmol, 9%): TLC R_f 0.29 (20:80 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$): δ 7.50 (1H, d, J = 1.8 Hz), 7.33 (1H, d, J = 8.0 Hz), 7.21 (1H, dd, J = 8.0, 1.8 Hz), 4.23 (1H, q, J = 7.3 Hz), 3.22 (3H, s), 3.19 (4H, br), 2.92 (1H, heptet, J = 7.0 Hz), 2.30 (3H, s), 1.55 (2H, br), 1.43-1.00 (8H, br), 1.27 (6H, d, J = 7.0 Hz), 0.91 (3H, br t, J = 7 Hz). MS (NH_3 -CI): m/e 492 (3), 491 (28), 490 (96), 489 (30), 488 (100). The second compound off the column was the title compound of Example 143 (1.30 g, 2.74 mmol, 80%): TLC R_f 0.22 (20:80 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$): δ 7.48 (1H, s), 7.21 (2H, s), 4.19 (2H, br), 3.21 (3H, s), 3.18 (4H, v br), 2.91 (1H, heptet, J = 6.6 Hz), 2.30 (3H, s), 1.55-1.46 (2H, m), 1.35-1.25 (2H, m), 1.27 (6H, d, J = 6.6 Hz), 1.11 (3H, br t, J = 7.0 Hz).

0.90 (3H, t, J = 7.0 Hz). MS (NH₃-CI): m/e 478 (3), 477 (28), 476 (98), 475 (30), 474 (100).

Example 24

5 Preparation of 8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine

A solution of the compound of Example 1 (650 mg, 1.41 mmol) in anhydrous tetrahydrofuran (5 mL) was treated with a tetrahydrofuran solution of borane (3 mL, 1 M, 3 mmol). The resulting solution was allowed to stir for 20 h, then was delivered slowly to stirring aq. 1 N sodium bicarbonate (10 mL). The mixture was stirred until gas evolution was complete, then was extracted with dichloromethane (twice 30 mL). The organic extracts were combined, dried over sodium sulfate, filtered and evaporated. The residual material was purified by elution through a short plug of silica gel (30:70 ethyl acetate-hexane), and evaporation gave the pure title product as an oil (429 mg, 1.04 mmol, 74%). Spectral data: TLC R_F 0.50 (30:70 ethyl acetate-hexane). ¹H NMR (300 MHz, CDCl₃): δ 7.50 (1H, d, J = 1.8 Hz), 7.24 (1H, d, J = 8.4 Hz), 7.18 (1H, dd, J = 8.4, 1.8 Hz), 3.88-3.39 (5H, br m), 3.22-3.09 (4H, m), 2.91 (1H, heptet, J = 7.0 Hz), 2.25 (3H, s), 1.52-1.41 (2H, m), 1.38-1.24 (2H, m), 1.27 (6H, d, J = 7.0 Hz), 1.08 (3H, t, J = 7.1 Hz), 0.91 (3H, t, J = 7.0 Hz). MS (ESI): m/e 450 (3), 449 (23), 448 (98), 447 (25), 446 (100). A solid derivative was obtained by precipitation of the hydrochloride salt from ether, melting point 79-81 °C.

Example 17

Preparation of 8-(2-bromo-4-isopropylphenyl)-2-methyl-4-(2-trifluoromethylphenyl)-5,6,7,8-tetrahydropteridin-6-one

35 **Part A.** A solution of 2-trifluoromethylphenylboronic acid (prepared according to the methods described in the review by N. Miyaura and A.

Suzuki, *Chem. Rev.* **1995**, 95, p. 2457) (1.00 g, 5.27 mmol), 4,6-dichloro-2-methyl-5-nitropyrimidine (0.91 g, 4.39 mmol) and tetrakis(triphenyl-phosphine) palladium (147 mg) in benzene (15 mL) was treated with aq. sodium carbonate solution (6 mL, 1 M). This mixture was heated to reflux for 6 h, then cooled and partitioned between water and ethyl acetate (60 mL each). The organic layer was washed with brine, and the aqueous layers were back-extracted in sequence with more ethyl acetate. The organic extracts were combined, dried over sodium sulfate, filtered and evaporated. The residue was separated by column chromatography (silica gel, 15:85 ethyl acetate-hexane) to afford 4-chloro-2-methyl-5-nitro-6-(2-trifluoromethylphenyl)pyrimidine (0.64 g, 2.01 mmol, 38%) as a waxy solid. Spectral data: R_f 0.40 (20:80 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$) δ 7.84-7.81 (1H, m), 7.69-7.63 (2H, m), 7.38-7.35 (1H, m), 2.84 (3H, s). MS (NH_3 -CI) m/e 320 (24), 319 (26), 318 (100).

Part B. A solution of the compound prepared in Part A above (1.07 g, 3.37 mmol) and 2-bromo-4-isopropylaniline (0.87 g, 4.04 mmol) in tetrahydrofuran (20 mL) was heated to reflux for 5 h, then cooled and poured into 100 mL ethyl acetate. This was washed with aq. sodium bicarbonate solution (100 mL, 1 N) and brine, then dried over sodium sulfate, filtered and evaporated. The resulting solid was triturated with 1:1 petroleum ether-diethyl ether, filtered and dried under vacuum to afford pure 4-(2-bromo-4-isopropylphenylamino)-2-methyl-5-nitro-6-(2-trifluoromethylphenyl)pyrimidine (1.51 g, 3.05 mmol, 90%). Spectral data: m.p. 152-154 °C. R_f 0.37 (20:80 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$) δ 10.20 (1H, br s), 8.26 (1H, d, J = 8.4 Hz), 7.78 (1H, d, J = 7.7 Hz), 7.65-7.57 (2H, m), 7.52 (1H, d, J = 1.8 Hz), 7.35 (1H, d, J = 7.0 Hz), 7.28 (1H, br s), 2.93 (1H, m), 2.66 (3H, s), 1.29 (6H, d, J = 7.0 Hz). MS (NH_3 -CI) m/e 498 (24), 497 (100), 496 (28), 495 (100).

Part C. The same procedure used for the reduction of a nitro group in Example 1, Part C was employed here. Thus, the compound of Part B above was converted to 5-amino-4-(2-bromo-4-isopropylphenylamino)-2-methyl-6-(2-trifluoromethylphenyl) pyrimidine in 32% yield. Spectral data: R_F 0.11 (20:80 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$) δ 8.58 (1H, d, $J = 8.4$ Hz), 7.83 (1H, d, $J = 7.7$ Hz), 7.67-7.58 (3H, m), 7.43 (1H, d, $J = 1.8$ Hz), 7.41 (1H, s), 7.25 (1H, dd, $J = 8.4, 1.8$ Hz), 2.89 (1H, m), 2.83 (2H, br s), 2.59 (3H, s), 1.26 (6H, d, $J = 7.0$ Hz). MS (NH_3 -CI) m/e 468 (24), 467 (100), 466 (30), 465 (99).

Part D. The same procedure used for the cyclization reaction of Example 1, Part D was employed here. Thus, the compound of Part C above was converted to the title compound in 60% yield. Spectral data: m.p. 238-239 °C. R_F 0.20 (50:50 ethyl acetate-hexane). 1H NMR (300 MHz, $CDCl_3$) δ 7.85 (1H, d, $J = 8.4$ Hz), 7.73-7.59 (3H, m), 7.48-7.41 (1H, m), 7.33 (1H, s), 7.30 (1H, s), 7.00 (1H, br s), 4.46 (1H, d, $J = 16.8$ Hz), 4.42 (1H, d, $J = 16.8$ Hz), 2.97 (1H, m), 2.40 (3H, s), 1.31 (6H, d, $J = 6.6$ Hz). MS (NH_3 -CI) m/e 508 (26), 507 (100), 506 (30), 505 (99).

25

Example 42

Preparation of Diethyl 2-[8-(2-bromo-4-isopropylphenyl)-2-methyl-6-oxo-5,6,7,8-tetrahydropteridin-4-yl]malonate

Sodium hydride suspension in mineral oil was washed with hexane and dried under vacuum, then taken up in tetrahydrofuran. This mixture is cooled to 0 °C, and treated with diethyl malonate (1.1 eq.). After the evolution of hydrogen gas is complete, the resulting solution is treated with 8-(2-bromo-4-isopropylphenyl)-4-chloro-2-methyl-5,6,7,8-tetrahydropteridin-6-one (see Example SF445, Part A, below; 1.0 eq.). The solution is heated to reflux until thin-layer chromatography shows the consumption of the starting material is nearly

complete. The mixture is allowed to cool, and poured into saturated aqueous ammonium chloride solution. This is extracted twice with ethyl acetate, and the extracts are washed with brine, combined, dried over sodium sulfate, filtered and evaporated. The residue is separated by column chromatography to afford the title product.

Example 131

10 Preparation of 8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridin-6-thione

A solution of the compound of Example 1 (323 mg, 0.70 mmol) in toluene (10 mL) was treated with Lawesson's Reagent (170 mg, 0.42 mmol). The resulting mixture was heated to reflux for 6 h, then cooled and evaporated. The residue was separated by column chromatography (silica gel, 10:90 ethyl acetate-hexane) to afford the title product, which was purified by recrystallization from ethyl acetate-hexane (280 mg, 0.59 mmol, 84%). Spectral data: m.p. 148-149 °C (ether-hexane). TLC R_f 0.31 (20:80 ethyl acetate-hexane). ^1H NMR (300 MHz, CDCl_3): δ 9.01 (1H, br s), 7.51 (1H, s), 7.22 (2H, s), 4.77 (1H, br d, $J = 15.7$ Hz), 4.56 (1H, br d, $J = 15.7$ Hz), 3.40-3.22 (4H, m), 2.92 (1H, heptet, $J = 7.0$ Hz), 2.29 (3H, s), 1.62-1.52 (2H, m), 1.40-1.30 (2H, m), 1.28 (6H, d, $J = 7.0$ Hz), 1.18 (3H, t, $J = 7.0$ Hz), 0.95 (3H, t, $J = 7.3$ Hz). MS (NH_3 -CI): m/e 479 (26), 478 (100), 477 (30), 476 (95). Analysis calc'd for $\text{C}_{22}\text{H}_{30}\text{BrN}_5\text{S}$: C, 55.46; H, 6.36; N, 14.70; found: C, 55.54; H, 6.38; N, 14.37.

Example 248Preparation of 8-(2-bromo-4-isopropyl-phenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridin-6-imine

5 The method of Robba et al., *Tetrahedron Letters*
1992, 33, p. 2803-2804 may be used here. Thus, a
solution of the compound of Example 131 in
tetrahydrofuran (0.5 M) is warmed to 55 °C, and treated
with 1.5 equivalents of mercuric chloride. Then, ammonia
10 gas is bubbled in, and addition is continued until five
minutes after the appearance of precipitating mercuric
sulfide. The reaction mixture is allowed to stir for an
additional hour, then is cooled, filtered through celite
and evaporated. The residue is triturated with a small
15 amount of water, filtered and dried to afford the title
compound.

Modifications of the procedures presented below for
Examples 501, 698 and 704 may be used to prepare many of
20 the compounds listed in Table III.

Example 501Preparation of 10-(2-bromo-4-(1-methylethyl)phenyl)-4-butyl-9,10-dihydro-2-methyl-4H,8H-pyrazino[3,2,1-d]pteridine-5,8(6H)-dione

25 **Part A.** Sodium hydride (mineral oil dispersion, 48
mg, 1.00 mmol) was washed with hexane, dried under
vacuum, and taken up in dimethylformamide (5 mL). To
this was added with stirring a solution of 5-amino-4-(2-
30 bromo-4-isopropylphenylamino)-6-chloro-2-
methylpyrimidine (311 mg, 0.87 mmol) in
dimethylformamide (5 mL). After stirring for 30 min.,
the mixture was treated with ethyl bromoacetate (0.10
mL, 0.90 mmol) in one portion. The mixture was allowed
35 to stir for 18 h, then poured into water (100 mL). This
was extracted twice with ethyl acetate (100 mL each),
and the organic extracts were washed in sequence with

two portions of water (100 mL each) and brine (100 mL), then combined, dried over sodium sulfate, filtered, and evaporated. The residual material was separated by column chromatography (silica gel, 20:80 ethyl acetate-hexane) to afford two fractions, the first being ethyl (8-(2-bromo-4-isopropylphenyl)-4-chloro-2-methyl-6-oxo-5,6,7,8-tetrahydropteridin-5-yl)acetate (89 mg, 0.18 mmol, 21%), the second being 8-(2-bromo-4-isopropylphenyl)-4-chloro-2-methyl-5,6,7,8-tetrahydropteridin-6-one (89 mg, 0.22 mmol, 26%). Spectral data for the first fraction: TLC R_f 0.25 (30:70 ethyl acetate-hexane). ^1H NMR (300 MHz, CDCl_3): δ 7.53 (1H, d, $J = 1.8$ Hz), 7.26 (1H, dd, $J = 8.4, 1.8$ Hz), 7.22 (1H, d, $J = 8.4$ Hz), 4.88 (1H, d, $J = 15$ Hz), 4.84 (1H, d, $J = 15$ Hz), 4.47 (1H, d, $J = 15.4$ Hz), 4.27 (1H, d, $J = 15.4$ Hz), 4.25 (2H, q, $J = 7.3$ Hz), 2.95 (1H, heptet, $J = 7.0$ Hz), 2.38 (3H, s), 1.29 (3H, t, $J = 7.3$ Hz), 1.28 (6H, d, $J = 7.0$ Hz). MS ($\text{NH}_3\text{-CI}$): m/e 486 (6), 485 (26), 484 (23), 483 (100), 482 (18), 481 (76).

Part B. A solution of the first fraction from Part A above (89 mg, 0.18 mmol) in butylamine (1.0 mL, 10.1 mmol) was treated with glacial acetic acid (0.011 mL, 0.19 mmol), and the resulting solution was heated to reflux for 20 h. The mixture was cooled and evaporated, and the residue was separated by column chromatography (silica gel, 15:85 ethyl acetate-hexane) to afford the title compound as a solid (50 mg, 0.10 mmol, 57%). Spectral data: m.p. 207-208 °C. TLC R_f 0.29 (30:70 ethyl acetate-hexane). ^1H NMR (300 MHz, CDCl_3): δ 7.54 (1H, d, $J = 1.8$ Hz), 7.27 (1H, dd, $J = 8.4, 1.8$ Hz), 7.23 (1H, d, $J = 8.4$ Hz), 4.66 (1H, d, $J = 18.6$ Hz), 4.55 (2H, br d, $J = 17$ Hz), 4.32 (1H, d, $J = 16.5$ Hz), 4.12 (2H, t, $J = 7.5$ Hz), 2.95 (1H, heptet, $J = 7.0$ Hz), 2.33 (3H, s), 1.70-1.59 (2H, m), 1.45-1.35 (2H, m), 1.29 (6H, d, $J = 7.0$ Hz), 0.96 (3H, t, $J = 7.1$ Hz). MS ($\text{NH}_3\text{-CI}$): m/e 476 (4), 475 (25), 474 (100), 473 (27), 472 (99). Analysis

calc'd for $C_{22}H_{26}BrN_5O_2$: C, 55.94; H, 5.56; N, 14.83;
found: C, 54.13; H, 5.34; N, 13.98.

5

Example 698Preparation of Diethyl 4-[2-bromo-4-(1-methylethyl)phenyl]-5,6,9,10-tetrahydro-2-methyl-6-oxo-4H,8H-pyrido[3,2,1-de]pteridine-10,10-dicarboxylate

Sodium hydride dispersion is washed free of oil
10 with hexane and dried under vacuum. Dimethylformamide is
added, and the mixture is cooled to 0 °C. A solution of
the compound of Example 42 (0.45 eq. based on the sodium
hydride) in dimethylformamide is added, and the mixture
15 is stirred for one hour. Then, a solution of 1,2-
dibromoethane (1.0 eq. based on the substrate) in
dimethylformamide is added dropwise slowly. The mixture
is allowed to stir overnight, then is poured into water.
The resulting mixture is extracted twice with ethyl
20 acetate, and the extracts are twice more washed with
water in sequence, then brine, then combined, dried over
sodium sulfate, filtered and evaporated. The residue is
separated by column chromatography to give the title
product.

25

Example 704Preparation of 4-[2-bromo-4-(1-methylethyl)phenyl]-5,6,9,10-tetrahydro-10,10-bis(methoxymethyl)-2-methyl-4H,8H-pyrido[3,2,1-de]pteridine

Part A. A solution of the compound of Example 698
30 in methylene chloride is cooled to 0 °C, and a solution
of diisobutylaluminum hydride (7 eq.) in methylene
chloride is slowly added by syringe. The mixture is
allowed to stir and warm to ambient temperature until
the substrate is consumed, then the reaction mixture is
35 slowly delivered to stirring ice-cold 1 N HCl solution.
The mixture is then neutralized to pH 7 with solid
sodium bicarbonate, and extracted twice with methylene

chloride. The extracts are combined, dried over sodium sulfate, filtered and evaporated. The residue is separated by column chromatography to afford 4-[2-bromo-4-(1-methylethyl)phenyl]-5,6,9,10-tetrahydro-10,10-bis(hydroxymethyl)-2-methyl-4*H*,8*H*-pyrido[3,2,1-*de*]pteridine.

Part B. Sodium hydride dispersion is washed free of oil with hexane, and dried under vacuum. Dimethylformamide is added, the suspension is cooled to 0 °C, and a solution of the compound of Part A above in dimethylformamide is slowly added. After stirring for 1 h, the mixture is treated with 2 eq. methyl iodide. The mixture is allowed to stir for at least 6 h, then poured into water. This is extracted twice with ethyl acetate, and the extracts are washed in sequence twice with water and once with brine, then combined, dried over sodium sulfate, filtered and evaporated. Chromatography is then used to isolate the title product.

Modifications of the procedures presented below for Examples 719 and 759 may be used to prepare many of the compounds listed in Table IV.

Example 719

Preparation of 5-[2-bromo-4-(1-methylethyl)phenyl]-*N*-butyl-*N*-ethyl-4,5-dihydro-7-methyltetrazolo[1,5-*f*]pteridin-9-amine

A solution of the compound of Example 1 (358 mg, 0.78 mmol) in tetrahydrofuran (10 mL) was treated with trimethylsilylazide (0.21 mL, 1.58 mmol), diethyl azodicarboxylate (0.25 mL, 1.59 mmol) and triphenylphosphine (408 mg, 1.56 mmol). The resulting solution was stirred for 20 h, then evaporated. The residual material was separated by column chromatography (silica gel, 20:80 ethyl acetate-hexane) to afford the title product as a low-melting solid (147 mg, 0.30 mmol, 39%). Spectral data: TLC *R_f* 0.37 (30:70 ethyl acetate-

hexane). ^1H NMR (300 MHz, CDCl_3): δ 7.52 (1H, d, $J = 1.8$ Hz), 7.26 (1H, dd, $J = 8.0, 1.8$ Hz), 7.21 (1H, d, $J = 8.0$ Hz), 5.23 (1H, d, $J = 15.0$ Hz), 4.98 (1H, d, $J = 15.0$ Hz), 3.56-3.48 (4H, m), 2.94 (1H, heptet, $J = 7.0$ Hz), 2.28 (3H, s), 1.69-1.59 (2H, m), 1.35-1.25 (2H, m), 1.29 (6H, d, $J = 7.0$ Hz), 1.21 (3H, t, $J = 7.1$ Hz), 0.90 (3H, t, $J = 7.3$ Hz). MS (NH_3 -CI): m/e 488 (17), 487 (64), 486 (18), 485 (63), 459 (100), 457 (97).

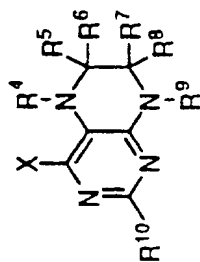
10

Example 759Preparation of 5-[2-bromo-4-(1-methylethyl)phenyl]-N-butyl-N-ethyl-5,6-dihydro-3-methylimidazo(1,2-f)pteridin-1-amine

A solution of the compound of Example 248 and a slight excess of bromoacetaldehyde diethyl acetal in ethanol is heated to reflux until the starting material is consumed. The reaction mixture is evaporated, and the residual material is purified by chromatography or recrystallization to afford the title product.

20

Table I.



Ex No	X	R ⁴	R ⁵	R ⁶	R ⁷	R ⁸	R ^{9 a}	R ¹⁰	mp, °C
1	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	H	H	A	CH ₃	146-147
2	(CH ₃) ₂ N	H	=O	---	H	H	A	CH ₃	---
3	(C ₂ H ₅) ₂ N	H	=O	---	H	H	A	CH ₃	---
4	(C ₄ H ₉) ₂ N	H	=O	---	H	H	A	CH ₃	---
5	CH ₃ OCH ₂ CH (C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	197-198
6	(CH ₃ O (CH ₂) ₂) ₂ N	H	=O	---	H	H	A	CH ₃	v. oil ^t
7	(CH ₃) ₂ NCH ₂ CH (C ₃ H ₇)NH	H	=O	---	H	H	A	CH ₃	188-189
8	C ₆ H ₅ CH ₂ (C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	---
9	(<i>c</i> -C ₃ H ₇)CH ₂ (C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	---
10	(C ₂ H ₅) ₂ CHNH	H	=O	---	H	H	A	CH ₃	---
11	C ₄ H ₉	H	=O	---	H	H	A	CH ₃	---
12	CH ₃	H	=O	---	H	H	A	CH ₃	---
13	(C ₂ H ₅) ₂ CH	H	=O	---	H	H	A	CH ₃	---
14	C ₆ H ₅	H	=O	---	H	H	A	CH ₃	---
15	<i>o</i> -ClC ₆ H ₄	H	=O	---	H	H	A	CH ₃	---
16	<i>o</i> -NO ₂ -C ₆ H ₄	H	=O	---	H	H	A	CH ₃	---
17	<i>o</i> -CF ₃ -C ₆ H ₄	H	=O	---	H	H	A	CH ₃	238-239

18	2-C ₅ H ₄ N	H	=O	---	H	H	A	CH ₃	---
19	(C ₂ H ₅ OC(=O)) ₂ CH	H	=O	---	H	H	A	CH ₃	---
20	(CN) ₂ CH	H	=O	---	H	H	A	CH ₃	---
21	(CH ₃ OCH ₂) ₂ CH	H	=O	---	H	H	A	CH ₃	---
22	CH ₃ S	H	=O	---	H	H	A	CH ₃	---
23	(C ₂ H ₅) ₂ CHO	H	=O	---	H	H	A	CH ₃	---
24	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	A	CH ₃	79-81 ^c
25	(CH ₃) ₂ N	H	H	H	H	H	A	CH ₃	---
26	(C ₂ H ₅) ₂ N	H	H	H	H	H	A	CH ₃	---
27	(C ₄ H ₉) ₂ N	H	H	H	H	H	A	CH ₃	---
28	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	140-145
29	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	A	CH ₃	v. oil ^d
30	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	A	CH ₃	83-85
31	C ₆ H ₅ CH ₂ (C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	---
32	(<i>c</i> -C ₃ H ₇)CH ₂ (C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	---
33	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	A	CH ₃	---
34	C ₄ H ₉ NH	H	H	H	H	H	A	CH ₃	160-162
35	CH ₃	H	H	H	H	H	A	CH ₃	---
36	(C ₂ H ₅) ₂ CH	H	H	H	H	H	A	CH ₃	---
37	C ₆ H ₅	H	H	H	H	H	A	CH ₃	---
38	<i>o</i> -ClC ₆ H ₄	H	H	H	H	H	A	CH ₃	---
39	<i>o</i> -NO ₂ -C ₆ H ₄	H	H	H	H	H	A	CH ₃	---
40	<i>o</i> -CF ₃ -C ₆ H ₄	H	H	H	H	H	A	CH ₃	108-110
41	2-C ₅ H ₄ N	H	H	H	H	H	A	CH ₃	---
42	(C ₂ H ₅ OC(=O)) ₂ CH	H	H	H	H	H	A	CH ₃	---

43	(CN) ₂ CH	H	H	H	H	A	CH ₃	---
44	(CH ₃ OCH ₂) ₂ CH	H	H	H	H	A	CH ₃	---
45	CH ₃ S	H	H	H	H	A	CH ₃	---
46	(C ₂ H ₅) ₂ CHO	H	H	H	H	A	CH ₃	---
47	C ₄ H ₉ (C ₂ H ₅)N	=O	---	---	---	B	CH ₃	171-173
48	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	=O	---	---	---	B	CH ₃	---
49	(CH ₃ O(CH ₂) ₂) ₂ N	=O	---	---	---	B	CH ₃	---
50	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	=O	---	---	---	B	CH ₃	---
51	O-CF ₃ -C ₆ H ₄	=O	---	---	---	B	CH ₃	---
52	(C ₂ H ₅) ₂ CHNH	=O	---	---	---	B	CH ₃	---
53	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	B	CH ₃	99-100
54	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	B	CH ₃	---
55	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	B	CH ₃	---
56	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	B	CH ₃	---
57	O-CF ₃ -C ₆ H ₄	H	H	H	H	B	CH ₃	---
58	(C ₂ H ₅) ₂ CHNH	H	H	H	H	B	CH ₃	---
59	C ₄ H ₉ (C ₂ H ₅)N	=O	---	---	---	C	CH ₃	157-159
60	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	=O	---	---	---	C	CH ₃	---
61	(CH ₃ O(CH ₂) ₂) ₂ N	=O	---	---	---	C	CH ₃	v. oil ^e
62	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	=O	---	---	---	C	CH ₃	---
63	O-CF ₃ -C ₆ H ₄	=O	---	---	---	C	CH ₃	---
64	(C ₂ H ₅) ₂ CHNH	=O	---	---	---	C	CH ₃	---
65	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	C	CH ₃	v. oil ^f
66	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	C	CH ₃	---
67	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	C	CH ₃	v. oil ^g

68	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	C	CH_3	---
69	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	C	CH_3	---
70	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	C	CH_3	---
71	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	---	D	CH_3	v. oil ^h
72	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	---	D	CH_3	209-210
73	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	---	D	CH_3	---
74	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	---	D	CH_3	---
75	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	---	D	CH_3	---
76	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	---	D	CH_3	---
77	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	D	CH_3	---
78	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	D	CH_3	---
79	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	D	CH_3	---
80	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	D	CH_3	---
81	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	D	CH_3	---
82	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	D	CH_3	---
83	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	---	E	CH_3	---
84	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	---	E	CH_3	107-109
85	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	---	E	CH_3	---
86	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	---	E	CH_3	---
87	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	---	E	CH_3	---
88	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	---	E	CH_3	---
89	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	E	CH_3	---
90	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	E	CH_3	---
91	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	E	CH_3	---
92	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	E	CH_3	---

93	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	CH ₃	---
94	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	CH ₃	---
95	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	---	---	CH ₃	---
96	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	---	---	CH ₃	227-229
97	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	---	---	CH ₃	---
98	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	---	---	CH ₃	---
99	O-CF ₃ -C ₆ H ₄	H	=O	---	---	---	CH ₃	---
100	(C ₂ H ₅) ₂ CHNH	H	=O	---	---	---	CH ₃	---
101	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	CH ₃	---
102	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	CH ₃	102-103
103	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	CH ₃	---
104	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	CH ₃	---
105	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	CH ₃	---
106	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	CH ₃	---
107	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	---	---	CH ₃	---
108	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	---	---	CH ₃	---
109	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	---	---	CH ₃	---
110	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	---	---	CH ₃	---
111	O-CF ₃ -C ₆ H ₄	H	=O	---	---	---	CH ₃	---
112	(C ₂ H ₅) ₂ CHNH	H	=O	---	---	---	CH ₃	---
113	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	CH ₃	---
114	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	CH ₃	---
115	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	CH ₃	---
116	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	CH ₃	---
117	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	CH ₃	---

118	$(C_2H_5)_2CHNH$	H	H	H	H	CH ₃	G	H	---
119	$C_4H_9(C_2H_5)N$	H	=O	---	---	CH ₃	J	H	---
120	$CH_3OCH_2CH(C_2H_5)NH$	H	=O	---	---	CH ₃	J	H	---
121	$(CH_3O(CH_2)_2)_2N$	H	=O	---	---	CH ₃	J	H	---
122	$(CH_3)_2NCH_2CH(C_3H_7)NH$	H	=O	---	---	CH ₃	J	H	---
123	$O-CF_3-C_6H_4$	H	=O	---	---	CH ₃	J	H	---
124	$(C_2H_5)_2CHNH$	H	=O	---	---	CH ₃	J	H	---
125	$C_4H_9(C_2H_5)N$	H	H	H	H	CH ₃	J	H	---
126	$CH_3OCH_2CH(C_2H_5)NH$	H	H	H	H	CH ₃	J	H	---
127	$(CH_3O(CH_2)_2)_2N$	H	H	H	H	CH ₃	J	H	---
128	$(CH_3)_2NCH_2CH(C_3H_7)NH$	H	H	H	H	CH ₃	J	H	---
129	$O-CF_3-C_6H_4$	H	H	H	H	CH ₃	J	H	---
130	$(C_2H_5)_2CHNH$	H	H	H	H	CH ₃	J	H	---
131	$C_4H_9(C_2H_5)N$	H	=S	---	---	CH ₃	A	H	148-149
132	$CH_3OCH_2CH(C_2H_5)NH$	H	=S	---	---	CH ₃	A	H	---
133	$(CH_3O(CH_2)_2)_2N$	H	=S	---	---	CH ₃	A	H	---
134	$(CH_3)_2NCH_2CH(C_3H_7)NH$	H	=S	---	---	CH ₃	A	H	---
135	$O-CF_3-C_6H_4$	H	=S	---	---	CH ₃	A	H	---
136	$(C_2H_5)_2CHNH$	H	=S	---	---	CH ₃	A	H	---
137	$C_4H_9(C_2H_5)N$	CH ₃	=S	---	---	CH ₃	A	H	v. oil
138	$CH_3OCH_2CH(C_2H_5)NH$	CH ₃	=S	---	---	CH ₃	A	H	---
139	$(CH_3O(CH_2)_2)_2N$	CH ₃	=S	---	---	CH ₃	A	H	---
140	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH ₃	=S	---	---	CH ₃	A	H	---
141	$O-CF_3-C_6H_4$	CH ₃	=S	---	---	CH ₃	A	H	---
142	$(C_2H_5)_2CHNH$	CH ₃	=S	---	---	CH ₃	A	H	---

143	$C_4H_9(C_2H_5)N$	CH_3	$=O$	---	H	H	A	CH_3	v. oil ^j
144	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	$=O$	---	H	H	A	CH_3	---
145	$(CH_3O(CH_2)_2)_2N$	CH_3	$=O$	---	H	H	A	CH_3	---
146	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	$=O$	---	H	H	A	CH_3	---
147	$O-CF_3-C_6H_4$	CH_3	$=O$	---	H	H	A	CH_3	v. oil ^k
148	$(C_2H_5)_2CHNH$	CH_3	$=O$	---	H	H	A	CH_3	---
149	$C_4H_9(C_2H_5)N$	CH_3	H	H	H	H	A	CH_3	v. oil ^l
150	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	H	H	H	H	A	CH_3	---
151	$(CH_3O(CH_2)_2)_2N$	CH_3	H	H	H	H	A	CH_3	---
152	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	H	H	H	H	A	CH_3	---
153	$O-CF_3-C_6H_4$	CH_3	H	H	H	H	A	CH_3	108-110
154	$(C_2H_5)_2CHNH$	CH_3	H	H	H	H	A	CH_3	---
155	$C_4H_9(C_2H_5)N$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	v. oil ^m
156	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	---
157	$(CH_3O(CH_2)_2)_2N$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	---
158	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	---
159	$O-CF_3-C_6H_4$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	v. oil ⁿ
160	$(C_2H_5)_2CHNH$	CH_3	$=O$	---	CH_3	CH_3	A	CH_3	---
161	$C_4H_9(C_2H_5)N$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
162	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
163	$(CH_3O(CH_2)_2)_2N$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
164	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
165	$O-CF_3-C_6H_4$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
166	$(C_2H_5)_2CHNH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
167	$C_4H_9(C_2H_5)N$	$CH_2CH=CH_2$	$=O$	---	H	H	A	CH_3	---

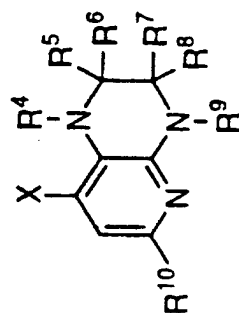
168	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
169	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
170	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
171	o-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	167-168
172	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
173	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
174	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
175	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
176	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
177	o-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
178	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
179	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
180	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
181	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
182	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
183	o-CF ₃ -C ₆ H ₄	CH ₃	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
184	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
185	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
186	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
187	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
188	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
189	o-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
190	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
191	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
192	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	CH ₃	CH ₃	H	H	A	CH ₃	---

193	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	CH_3	CH_3	H	H	A	CH_3	---
194	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	CH_3	CH_3	H	H	A	CH_3	---
195	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	CH_3	CH_3	H	H	A	CH_3	---
196	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	CH_3	CH_3	H	H	A	CH_3	---
197	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	CH_3	CH_3	A	CH_3	---
198	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
199	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	CH_3	CH_3	A	CH_3	---
200	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
201	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	CH_3	CH_3	A	CH_3	---
202	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
203	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	CH_3	CH_3	A	CH_3	---
204	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	CH_3	CH_3	A	CH_3	---
205	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	CH_3	CH_3	A	CH_3	---
206	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	CH_3	CH_3	A	CH_3	---
207	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	CH_3	CH_3	A	CH_3	---
208	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	CH_3	CH_3	A	CH_3	---
209	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	H	H	A	H	---
210	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	H	H	A	H	---
211	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	H	H	A	H	---
212	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	H	H	A	H	---
213	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	H	H	A	H	---
214	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	H	H	A	H	---
215	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	H	A	H	---
216	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	H	A	H	---
217	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	H	A	H	---

218	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	A	H	---
219	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	A	H	---
220	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	A	H	---
221	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	---	A	CN	---
222	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	---	A	CN	---
223	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	---	A	CN	---
224	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	---	A	CN	---
225	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	---	A	CN	---
226	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	---	A	CN	---
227	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	A	CN	---
228	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	A	CN	---
229	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	A	CN	---
230	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	A	CN	---
231	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	A	CN	---
232	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	A	CN	---
233	2-morpholinyl	H	H	H	H	A	CH ₃	60-62
234	2-morpholinyl	H	=O	---	---	B	CH ₃	---
235	2-morpholinyl	H	=O	---	---	C	CH ₃	---
236	2-morpholinyl	H	=O	---	---	D	CH ₃	---
237	2-morpholinyl	H	=O	---	---	E	CH ₃	---
238	2-morpholinyl	H	=O	---	---	F	CH ₃	---
239	2-morpholinyl	H	=O	---	---	G	CH ₃	---
240	2-morpholinyl	H	=O	---	---	J	CH ₃	---
241	2-morpholinyl	H	H	H	H	B	CH ₃	---
242	2-morpholinyl	H	H	H	H	C	CH ₃	---

243	2-morpholinyl	H	H	H	H	H	D	CH ₃	---
244	2-morpholinyl	H	H	H	H	H	E	CH ₃	---
245	2-morpholinyl	H	H	H	H	H	F	CH ₃	---
246	2-morpholinyl	H	H	H	H	H	G	CH ₃	---
247	2-morpholinyl	H	H	H	H	H	J	CH ₃	---
248	C ₄ H ₉ (C ₂ H ₅)N	H	=NH	---	H	H	A	CH ₃	---
249	C ₄ H ₉ (C ₂ H ₅)N	H	=NH	---	H	H	C	CH ₃	---
250	C ₄ H ₉ (C ₂ H ₅)N	H	=NC ₄ H ₉	---	H	H	A	CH ₃	---

Table II.



Ex No	X	R ⁴	R ⁵	R ⁶	R ⁷	R ⁸	R ^{9 a}	R ¹⁰	mp, °C
251	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	H	H	A	CH ₃	---
252	(CH ₃) ₂ N	H	=O	---	H	H	A	CH ₃	---
253	(C ₂ H ₅) ₂ N	H	=O	---	H	H	A	CH ₃	---
254	(C ₄ H ₉) ₂ N	H	=O	---	H	H	A	CH ₃	---
255	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	---
256	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	H	H	A	CH ₃	---
257	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	H	H	A	CH ₃	---
258	C ₆ H ₅ CH ₂ (C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	---
259	(C-C ₃ H ₇)CH ₂ (C ₂ H ₅)NH	H	=O	---	H	H	A	CH ₃	---
260	(C ₂ H ₅) ₂ CHNH	H	=O	---	H	H	A	CH ₃	---
261	C ₄ H ₉	H	=O	---	H	H	A	CH ₃	---
262	CH ₃	H	=O	---	H	H	A	CH ₃	---
263	(C ₂ H ₅) ₂ CH	H	=O	---	H	H	A	CH ₃	---
264	C ₆ H ₅	H	=O	---	H	H	A	CH ₃	---
265	O-ClC ₆ H ₄	H	=O	---	H	H	A	CH ₃	---
266	O-NO ₂ -C ₆ H ₄	H	=O	---	H	H	A	CH ₃	---
267	O-CF ₃ -C ₆ H ₄	H	=O	---	H	H	A	CH ₃	---

268	2-C ₅ H ₄ N	H	=O	---	H	H	A	CH ₃	---
269	(C ₂ H ₅ OC(=O)) ₂ CH	H	=O	---	H	H	A	CH ₃	---
270	(CN) ₂ CH	H	=O	---	H	H	A	CH ₃	---
271	(CH ₃ OCH ₂) ₂ CH	H	=O	---	H	H	A	CH ₃	---
272	CH ₃ S	H	=O	---	H	H	A	CH ₃	---
273	(C ₂ H ₅) ₂ CHO	H	=O	---	H	H	A	CH ₃	---
274	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	A	CH ₃	---
275	(CH ₃) ₂ N	H	H	H	H	H	A	CH ₃	---
276	(C ₂ H ₅) ₂ N	H	H	H	H	H	A	CH ₃	---
277	(C ₄ H ₉) ₂ N	H	H	H	H	H	A	CH ₃	---
278	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	---
279	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	A	CH ₃	---
280	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	A	CH ₃	---
281	C ₆ H ₅ CH ₂ (C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	---
282	(C-C ₃ H ₇)CH ₂ (C ₂ H ₅)NH	H	H	H	H	H	A	CH ₃	---
283	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	A	CH ₃	---
284	C ₄ H ₉ NH	H	H	H	H	H	A	CH ₃	---
285	CH ₃	H	H	H	H	H	A	CH ₃	---
286	(C ₂ H ₅) ₂ CH	H	H	H	H	H	A	CH ₃	---
287	C ₅ H ₅	H	H	H	H	H	A	CH ₃	---
288	O-ClC ₆ H ₄	H	H	H	H	H	A	CH ₃	---
289	O-NO ₂ -C ₆ H ₄	H	H	H	H	H	A	CH ₃	---
290	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	A	CH ₃	---
291	2-C ₅ H ₄ N	H	H	H	H	H	A	CH ₃	---
292	(C ₂ H ₅ OC(=O)) ₂ CH	H	H	H	H	H	A	CH ₃	---

293	(CN) ₂ CH	H	H	H	H	A	CH ₃	---
294	(CH ₃ OCH ₂) ₂ CH	H	H	H	H	A	CH ₃	---
295	CH ₃ S	H	H	H	H	A	CH ₃	---
296	(C ₂ H ₅) ₂ CHO	H	H	H	H	A	CH ₃	---
297	C ₄ H ₉ (C ₂ H ₅)N	=O	---	---	---	B	CH ₃	---
298	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	=O	---	---	---	B	CH ₃	---
299	(CH ₃ O(CH ₂) ₂) ₂ N	=O	---	---	---	B	CH ₃	---
300	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	=O	---	---	---	B	CH ₃	---
301	O-CF ₃ -C ₆ H ₄	=O	---	---	---	B	CH ₃	---
302	(C ₂ H ₅) ₂ CHNH	=O	---	---	---	B	CH ₃	---
303	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	B	CH ₃	---
304	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	B	CH ₃	---
305	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	B	CH ₃	---
306	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	B	CH ₃	---
307	O-CF ₃ -C ₆ H ₄	H	H	H	H	B	CH ₃	---
308	(C ₂ H ₅) ₂ CHNH	H	H	H	H	B	CH ₃	---
309	C ₄ H ₉ (C ₂ H ₅)N	=O	---	---	---	C	CH ₃	---
310	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	=O	---	---	---	C	CH ₃	---
311	(CH ₃ O(CH ₂) ₂) ₂ N	=O	---	---	---	C	CH ₃	---
312	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	=O	---	---	---	C	CH ₃	---
313	O-CF ₃ -C ₆ H ₄	=O	---	---	---	C	CH ₃	---
314	(C ₂ H ₅) ₂ CHNH	=O	---	---	---	C	CH ₃	---
315	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	C	CH ₃	---
316	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	C	CH ₃	---
317	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	C	CH ₃	---

318	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	C	CH ₃	---
319	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	C	CH ₃	---
320	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	C	CH ₃	---
321	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	H	D	CH ₃	---
322	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	H	D	CH ₃	---
323	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	H	D	CH ₃	---
324	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	H	D	CH ₃	---
325	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	H	D	CH ₃	---
326	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	H	D	CH ₃	---
327	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	D	CH ₃	---
328	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	D	CH ₃	---
329	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	D	CH ₃	---
330	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	D	CH ₃	---
331	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	D	CH ₃	---
332	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	D	CH ₃	---
333	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	H	E	CH ₃	---
334	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	H	E	CH ₃	---
335	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	H	E	CH ₃	---
336	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	H	E	CH ₃	---
337	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	H	E	CH ₃	---
338	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	H	E	CH ₃	---
339	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	E	CH ₃	---
340	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	E	CH ₃	---
341	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	E	CH ₃	---
342	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	E	CH ₃	---

343	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	H	H	E	CH ₃	---
344	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	H	H	E	CH ₃	---
345	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	---	---	---	---	F	CH ₃	---
346	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	---	---	---	---	F	CH ₃	---
347	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	---	---	---	---	F	CH ₃	---
348	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	---	---	---	---	F	CH ₃	---
349	O-CF ₃ -C ₆ H ₄	H	=O	---	---	---	---	---	F	CH ₃	---
350	(C ₂ H ₅) ₂ CHNH	H	=O	---	---	---	---	---	F	CH ₃	---
351	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	H	H	F	CH ₃	---
352	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	H	H	F	CH ₃	---
353	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	H	H	F	CH ₃	---
354	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	H	H	F	CH ₃	---
355	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	H	H	F	CH ₃	---
356	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	H	H	F	CH ₃	---
357	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	---	---	---	---	G	CH ₃	---
358	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	---	---	---	---	G	CH ₃	---
359	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	---	---	---	---	G	CH ₃	---
360	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	---	---	---	---	G	CH ₃	---
361	O-CF ₃ -C ₆ H ₄	H	=O	---	---	---	---	---	G	CH ₃	---
362	(C ₂ H ₅) ₂ CHNH	H	=O	---	---	---	---	---	G	CH ₃	---
363	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	H	H	H	H	G	CH ₃	---
364	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	H	H	H	H	G	CH ₃	---
365	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	H	H	H	H	G	CH ₃	---
366	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	H	H	H	H	G	CH ₃	---
367	O-CF ₃ -C ₆ H ₄	H	H	H	H	H	H	H	G	CH ₃	---

368	(C ₂ H ₅) ₂ CHNH	H	H	H	H	H	H	CH ₃	G	H	---
369	C ₄ H ₉ (C ₂ H ₅)N	H	=O	---	---	---	---	CH ₃	J	H	---
370	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=O	---	---	---	---	CH ₃	J	H	---
371	(CH ₃ O(CH ₂) ₂) ₂ N	H	=O	---	---	---	---	CH ₃	J	H	---
372	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=O	---	---	---	---	CH ₃	J	H	---
373	O-CF ₃ -C ₆ H ₄	H	=O	---	---	---	---	CH ₃	J	H	---
374	(C ₂ H ₅) ₂ CHNH	H	=O	---	---	---	---	CH ₃	J	H	---
375	C ₄ H ₉ (C ₂ H ₅)N	H	H	H	---	---	---	CH ₃	J	H	---
376	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	H	H	---	---	---	CH ₃	J	H	---
377	(CH ₃ O(CH ₂) ₂) ₂ N	H	H	H	---	---	---	CH ₃	J	H	---
378	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	H	H	---	---	---	CH ₃	J	H	---
379	O-CF ₃ -C ₆ H ₄	H	H	H	---	---	---	CH ₃	J	H	---
380	(C ₂ H ₅) ₂ CHNH	H	H	H	---	---	---	CH ₃	J	H	---
381	C ₄ H ₉ (C ₂ H ₅)N	H	=S	---	---	---	---	CH ₃	A	H	---
382	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	=S	---	---	---	---	CH ₃	A	H	---
383	(CH ₃ O(CH ₂) ₂) ₂ N	H	=S	---	---	---	---	CH ₃	A	H	---
384	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	H	=S	---	---	---	---	CH ₃	A	H	---
385	O-CF ₃ -C ₆ H ₄	H	=S	---	---	---	---	CH ₃	A	H	---
386	(C ₂ H ₅) ₂ CHNH	H	=S	---	---	---	---	CH ₃	A	H	---
387	C ₄ H ₉ (C ₂ H ₅)N	CH ₃	=S	---	---	---	---	CH ₃	A	H	---
388	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₃	=S	---	---	---	---	CH ₃	A	H	---
389	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₃	=S	---	---	---	---	CH ₃	A	H	---
390	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₃	=S	---	---	---	---	CH ₃	A	H	---
391	O-CF ₃ -C ₆ H ₄	CH ₃	=S	---	---	---	---	CH ₃	A	H	---
392	(C ₂ H ₅) ₂ CHNH	CH ₃	=S	---	---	---	---	CH ₃	A	H	---

393	$C_4H_9(C_2H_5)N$	CH_3	=O	---	H	H	A	CH_3	---
394	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	=O	---	H	H	A	CH_3	---
395	$(CH_3O(CH_2)_2)_2N$	CH_3	=O	---	H	H	A	CH_3	---
396	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	=O	---	H	H	A	CH_3	---
397	$O-CF_3-C_6H_4$	CH_3	=O	---	H	H	A	CH_3	---
398	$(C_2H_5)_2CHNH$	CH_3	=O	---	H	H	A	CH_3	---
399	$C_4H_9(C_2H_5)N$	CH_3	H	H	H	H	A	CH_3	---
400	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	H	H	H	H	A	CH_3	---
401	$(CH_3O(CH_2)_2)_2N$	CH_3	H	H	H	H	A	CH_3	---
402	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	H	H	H	H	A	CH_3	---
403	$O-CF_3-C_6H_4$	CH_3	H	H	H	H	A	CH_3	---
404	$(C_2H_5)_2CHNH$	CH_3	H	H	H	H	A	CH_3	---
405	$C_4H_9(C_2H_5)N$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
406	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
407	$(CH_3O(CH_2)_2)_2N$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
408	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
409	$O-CF_3-C_6H_4$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
410	$(C_2H_5)_2CHNH$	CH_3	=O	---	CH_3	CH_3	A	CH_3	---
411	$C_4H_9(C_2H_5)N$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
412	$CH_3OCH_2CH(C_2H_5)NH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
413	$(CH_3O(CH_2)_2)_2N$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
414	$(CH_3)_2NCH_2CH(C_3H_7)NH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
415	$O-CF_3-C_6H_4$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
416	$(C_2H_5)_2CHNH$	CH_3	H	H	CH_3	CH_3	A	CH_3	---
417	$C_4H_9(C_2H_5)N$	$CH_2CH=CH_2$	=O	---	H	H	A	CH_3	---

418	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
419	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
420	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
421	O-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
422	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	=O	---	H	H	A	CH ₃	---
423	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
424	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
425	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
426	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
427	O-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
428	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	H	H	H	H	A	CH ₃	---
429	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
430	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
431	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
432	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
433	O-CF ₃ -C ₆ H ₄	CH ₃	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
434	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	=O	---	CH ₂ CH=CH ₂	H	A	CH ₃	---
435	C ₄ H ₉ (C ₂ H ₅)N	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
436	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
437	(CH ₃ O(CH ₂) ₂) ₂ N	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
438	(CH ₃) ₂ NCH ₂ CH(C ₃ H ₇)NH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
439	O-CF ₃ -C ₆ H ₄	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
440	(C ₂ H ₅) ₂ CHNH	CH ₂ CH=CH ₂	H	H	CH ₂ CH=CH ₂	H	A	CH ₃	---
441	C ₄ H ₉ (C ₂ H ₅)N	H	CH ₃	CH ₃	H	H	A	CH ₃	---
442	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	H	CH ₃	CH ₃	H	H	A	CH ₃	---

443	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	CH_3	CH_3	H	H	A	CH_3	---
444	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	CH_3	CH_3	H	H	A	CH_3	---
445	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	CH_3	CH_3	H	H	A	CH_3	---
446	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	CH_3	CH_3	H	H	A	CH_3	---
447	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	CH_3	CH_3	A	CH_3	---
448	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
449	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	CH_3	CH_3	A	CH_3	---
450	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
451	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	CH_3	CH_3	A	CH_3	---
452	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	CH_3	CH_3	A	CH_3	---
453	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	CH_3	CH_3	A	CH_3	---
454	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	CH_3	CH_3	A	CH_3	---
455	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	CH_3	CH_3	A	CH_3	---
456	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	CH_3	CH_3	A	CH_3	---
457	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	CH_3	CH_3	A	CH_3	---
458	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	CH_3	CH_3	A	CH_3	---
459	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	H	H	A	H	---
460	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	H	H	A	H	---
461	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	H	H	A	H	---
462	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	H	H	A	H	---
463	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	H	H	A	H	---
464	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	H	H	A	H	---
465	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	H	A	H	---
466	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	H	A	H	---
467	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	H	A	H	---

468	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	A	H	---
469	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	A	H	---
470	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	A	H	---
471	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	=O	---	H	A	CN	---
472	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	=O	---	H	A	CN	---
473	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	=O	---	H	A	CN	---
474	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	=O	---	H	A	CN	---
475	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	=O	---	H	A	CN	---
476	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	=O	---	H	A	CN	---
477	$\text{C}_4\text{H}_9(\text{C}_2\text{H}_5)\text{N}$	H	H	H	H	A	CN	---
478	$\text{CH}_3\text{OCH}_2\text{CH}(\text{C}_2\text{H}_5)\text{NH}$	H	H	H	H	A	CN	---
479	$(\text{CH}_3\text{O}(\text{CH}_2)_2)_2\text{N}$	H	H	H	H	A	CN	---
480	$(\text{CH}_3)_2\text{NCH}_2\text{CH}(\text{C}_3\text{H}_7)\text{NH}$	H	H	H	H	A	CN	---
481	$\text{O}-\text{CF}_3-\text{C}_6\text{H}_4$	H	H	H	H	A	CN	---
482	$(\text{C}_2\text{H}_5)_2\text{CHNH}$	H	H	H	H	A	CN	---
483	2-morpholinyl	H	H	H	H	A	CH_3	---
484	2-morpholinyl	H	=O	---	H	B	CH_3	---
485	2-morpholinyl	H	=O	---	H	C	CH_3	---
486	2-morpholinyl	H	=O	---	H	D	CH_3	---
487	2-morpholinyl	H	=O	---	H	E	CH_3	---
488	2-morpholinyl	H	=O	---	H	F	CH_3	---
489	2-morpholinyl	H	=O	---	H	G	CH_3	---
490	2-morpholinyl	H	=O	---	H	J	CH_3	---
491	2-morpholinyl	H	H	H	H	B	CH_3	---
492	2-morpholinyl	H	H	H	H	C	CH_3	---

493	2-morpholinyl	H	H	H	H	H	D	CH ₃	---
494	2-morpholinyl	H	H	H	H	H	E	CH ₃	---
495	2-morpholinyl	H	H	H	H	H	F	CH ₃	---
496	2-morpholinyl	H	H	H	H	H	G	CH ₃	---
497	2-morpholinyl	H	H	H	H	H	J	CH ₃	---
498	C ₄ H ₉ (C ₂ H ₅)N	H	=NH	---	---	---	A	CH ₃	---
499	C ₄ H ₉ (C ₂ H ₅)N	H	=NH	---	---	---	C	CH ₃	---
500	C ₄ H ₉ (C ₂ H ₅)N	H	=NC ₄ H ₉	---	---	---	A	CH ₃	---

Key: (a) R⁹ group codes: A =

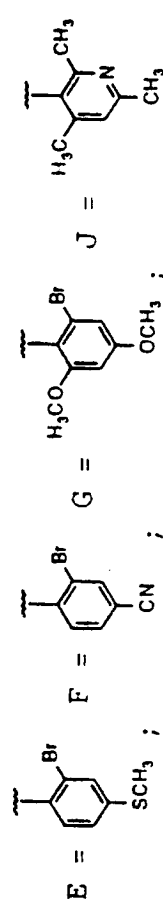
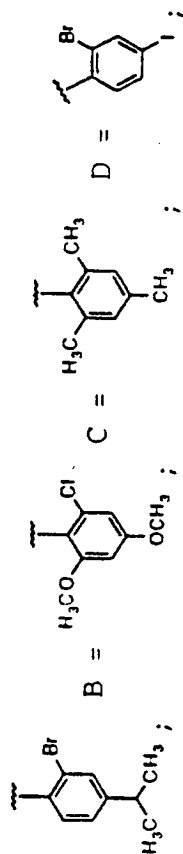
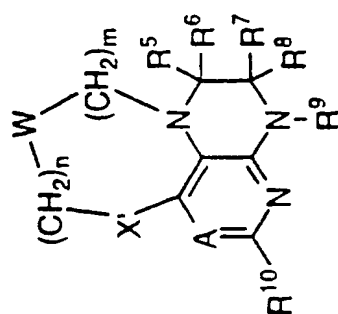


Table III.



Ex. No.	A	X'	n	W	m	R ⁵	R ⁶	R ⁷	R ⁸	R ^{9a}	R ¹⁰	m.p.
501	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	A	CH ₃	207-208
502	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	B	CH ₃	—
503	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	C	CH ₃	—
504	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	D	CH ₃	—
505	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	E	CH ₃	—
506	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	F	CH ₃	—
507	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	G	CH ₃	—
508	N	C ₄ H ₉ N	0	C=O	1	=O	—	H	H	J	CH ₃	—
509	N	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	—	H	H	A	CH ₃	—
510	N	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	—	H	H	B	CH ₃	—
511	N	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	—	H	H	C	CH ₃	—
512	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	A	CH ₃	—
513	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	B	CH ₃	—
514	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	C	CH ₃	—
515	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	A	CH ₃	—
516	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	B	CH ₃	—
517	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	—	H	H	C	CH ₃	—

518	N	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
519	N	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
520	N	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	C	CH ₃	---
521	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	A	CH ₃	---
522	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	B	CH ₃	---
523	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	C	CH ₃	---
524	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
525	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
526	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	C	CH ₃	---
527	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
528	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
529	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	C	CH ₃	---
530	N	C ₄ H ₉ N	1	CH ₂	1	=O	---	---	---	A	CH ₃	---
531	N	C ₄ H ₉ N	1	CH ₂	1	=O	---	---	---	B	CH ₃	---
532	N	C ₄ H ₉ N	1	CH ₂	1	=O	---	---	---	C	CH ₃	---
533	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	---	---	A	CH ₃	---
534	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	---	---	B	CH ₃	---
535	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	---	---	C	CH ₃	---
536	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	A	CH ₃	---
537	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	B	CH ₃	---
538	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	C	CH ₃	---
539	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	A	CH ₃	---
540	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	B	CH ₃	---
541	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	---	---	C	CH ₃	---
542	N	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
543	N	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
544	N	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	C	CH ₃	---

545	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	A	CH ₃	---
546	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	B	CH ₃	---
547	N	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	C	CH ₃	---
548	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
549	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
550	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	C	CH ₃	---
551	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
552	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
553	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	C	CH ₃	---
554	N	C ₄ H ₉ N	0	CH ₂	1	=O	---	---	---	A	CH ₃	---
555	N	C ₄ H ₉ N	0	CH ₂	1	=O	---	---	---	B	CH ₃	---
556	N	C ₄ H ₉ N	0	CH ₂	1	=O	---	---	---	C	CH ₃	---
557	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	---	---	A	CH ₃	---
558	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	---	---	B	CH ₃	---
559	N	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	---	---	C	CH ₃	---
560	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	A	CH ₃	---
561	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	B	CH ₃	---
562	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	C	CH ₃	---
563	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	A	CH ₃	---
564	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	B	CH ₃	---
565	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	---	---	C	CH ₃	---
566	N	C ₄ H ₉ N	1	C=O	1	=O	---	---	---	A	CH ₃	---
567	N	C ₄ H ₉ N	1	C=O	1	=O	---	---	---	B	CH ₃	---
568	N	C ₄ H ₉ N	1	C=O	1	=O	---	---	---	C	CH ₃	---
569	N	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	---	---	A	CH ₃	---
570	N	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	---	---	B	CH ₃	---
571	N	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	---	---	C	CH ₃	---

572	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	A	CH ₃	---
573	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	B	CH ₃	---
574	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	C	CH ₃	---
575	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	A	CH ₃	---
576	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	B	CH ₃	---
577	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	H	H	C	CH ₃	---
578	N	C ₄ H ₉ N	1	C=O	0	H	H	H	A	CH ₃	---
579	N	C ₄ H ₉ N	1	C=O	0	H	H	H	B	CH ₃	---
580	N	C ₄ H ₉ N	1	C=O	0	H	H	H	C	CH ₃	---
581	N	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	A	CH ₃	---
582	N	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	B	CH ₃	---
583	N	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	C	CH ₃	---
584	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	A	CH ₃	---
585	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	B	CH ₃	---
586	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	C	CH ₃	---
587	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	A	CH ₃	---
588	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	B	CH ₃	---
589	N	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	C	CH ₃	---
590	N	C ₄ H ₉ N	0	C=O	0	H	H	H	A	CH ₃	---
591	N	C ₄ H ₉ N	0	C=O	0	H	H	H	B	CH ₃	---
592	N	C ₄ H ₉ N	0	C=O	0	H	H	H	C	CH ₃	---
593	N	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	A	CH ₃	---
594	N	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	B	CH ₃	---
595	N	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	C	CH ₃	---
596	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	A	CH ₃	---
597	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	B	CH ₃	---
598	N	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	C	CH ₃	---

599	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	A	CH ₃	---
600	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	B	CH ₃	---
601	N	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	C	CH ₃	---
602	CH	C ₄ H ₉ N	0	C=O	1	=O	---	H	H	A	CH ₃	---
603	CH	C ₄ H ₉ N	0	C=O	1	=O	---	H	H	B	CH ₃	---
604	CH	C ₄ H ₉ N	0	C=O	1	=O	---	H	H	C	CH ₃	---
605	CH	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	---	H	H	A	CH ₃	---
606	CH	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	---	H	H	B	CH ₃	---
607	CH	(C ₂ H ₅) ₂ CHN	0	C=O	1	=O	---	H	H	C	CH ₃	---
608	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	A	CH ₃	---
609	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	B	CH ₃	---
610	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	C	CH ₃	---
611	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	A	CH ₃	---
612	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	B	CH ₃	---
613	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	1	=O	---	H	H	C	CH ₃	---
614	CH	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
615	CH	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
616	CH	C ₄ H ₉ N	0	CH ₂	1	H	H	H	H	C	CH ₃	---
617	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	A	CH ₃	---
618	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	B	CH ₃	---
619	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	H	H	H	H	C	CH ₃	---
620	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
621	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
622	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	C	CH ₃	---
623	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	A	CH ₃	---
624	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	B	CH ₃	---
625	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	H	H	H	H	C	CH ₃	---

626	CH	C ₄ H ₉ N	1	CH ₂	1	=O	---	H	H	A	CH ₃	---
627	CH	C ₄ H ₉ N	1	CH ₂	1	=O	---	H	H	B	CH ₃	---
628	CH	C ₄ H ₉ N	1	CH ₂	1	=O	---	H	H	C	CH ₃	---
629	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	H	H	A	CH ₃	---
630	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	H	H	B	CH ₃	---
631	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	=O	---	H	H	C	CH ₃	---
632	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	A	CH ₃	---
633	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	B	CH ₃	---
634	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	C	CH ₃	---
635	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	A	CH ₃	---
636	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	B	CH ₃	---
637	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	=O	---	H	H	C	CH ₃	---
638	CH	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
639	CH	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
640	CH	C ₄ H ₉ N	1	CH ₂	1	H	H	H	H	C	CH ₃	---
641	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	A	CH ₃	---
642	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	B	CH ₃	---
643	CH	(C ₂ H ₅) ₂ CHN	1	CH ₂	1	H	H	H	H	C	CH ₃	---
644	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
645	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
646	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	C	CH ₃	---
647	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	A	CH ₃	---
648	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	B	CH ₃	---
649	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	CH ₂	1	H	H	H	H	C	CH ₃	---
650	CH	C ₄ H ₉ N	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
651	CH	C ₄ H ₉ N	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
652	CH	C ₄ H ₉ N	0	CH ₂	1	=O	---	H	H	C	CH ₃	---

653	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
654	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
655	CH	(C ₂ H ₅) ₂ CHN	0	CH ₂	1	=O	---	H	H	C	CH ₃	---
656	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
657	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
658	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	C	CH ₃	---
659	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
660	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
661	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	CH ₂	1	=O	---	H	H	C	CH ₃	---
662	CH	C ₄ H ₉ N	1	C=O	1	=O	---	H	H	A	CH ₃	---
663	CH	C ₄ H ₉ N	1	C=O	1	=O	---	H	H	B	CH ₃	---
664	CH	C ₄ H ₉ N	1	C=O	1	=O	---	H	H	C	CH ₃	---
665	CH	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	H	H	A	CH ₃	---
666	CH	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	H	H	B	CH ₃	---
667	CH	(C ₂ H ₅) ₂ CHN	1	C=O	1	=O	---	H	H	C	CH ₃	---
668	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	A	CH ₃	---
669	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	B	CH ₃	---
670	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	C	CH ₃	---
671	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	A	CH ₃	---
672	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	B	CH ₃	---
673	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	1	=O	---	H	H	C	CH ₃	---
674	CH	C ₄ H ₉ N	1	C=O	0	H	H	H	H	A	CH ₃	---
675	CH	C ₄ H ₉ N	1	C=O	0	H	H	H	H	B	CH ₃	---
676	CH	C ₄ H ₉ N	1	C=O	0	H	H	H	H	C	CH ₃	---
677	CH	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	H	A	CH ₃	---
678	CH	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	H	B	CH ₃	---
679	CH	(C ₂ H ₅) ₂ CHN	1	C=O	0	H	H	H	H	C	CH ₃	---

680	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	A	CH ₃	---
681	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	B	CH ₃	---
682	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	C	CH ₃	---
683	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	A	CH ₃	---
684	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	B	CH ₃	---
685	CH	C ₄ H ₉ CH(C ₂ H ₅)N	1	C=O	0	H	H	H	H	C	CH ₃	---
686	CH	C ₄ H ₉	0	C=O	0	H	H	H	H	A	CH ₃	---
687	CH	C ₄ H ₉	0	C=O	0	H	H	H	H	B	CH ₃	---
688	CH	C ₄ H ₉	0	C=O	0	H	H	H	H	C	CH ₃	---
689	CH	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	H	A	CH ₃	---
690	CH	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	H	B	CH ₃	---
691	CH	(C ₂ H ₅) ₂ CHN	0	C=O	0	H	H	H	H	C	CH ₃	---
692	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	A	CH ₃	---
693	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	B	CH ₃	---
694	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	C	CH ₃	---
695	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	A	CH ₃	---
696	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	B	CH ₃	---
697	CH	C ₄ H ₉ CH(C ₂ H ₅)N	0	C=O	0	H	H	H	H	C	CH ₃	---
698	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
699	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
700	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	=O	---	H	H	C	CH ₃	---
701	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	H	H	H	H	A	CH ₃	---
702	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	H	H	H	H	B	CH ₃	---
703	N	[C ₂ H ₅ OC(=O)] ₂ C	0	CH ₂	1	H	H	H	H	C	CH ₃	---
704	N	(CH ₃ OCH ₂) ₂ C	0	CH ₂	1	H	H	H	H	A	CH ₃	---
705	N	(CH ₃ OCH ₂) ₂ C	0	CH ₂	1	H	H	H	H	B	CH ₃	---
706	N	(CH ₃ OCH ₂) ₂ C	0	CH ₂	1	H	H	H	H	C	CH ₃	---

707	N	(NC) ₂ C	0	CH ₂	1	=O	---	H	H	A	CH ₃	---
708	N	(NC) ₂ C	0	CH ₂	1	=O	---	H	H	B	CH ₃	---
709	N	(NC) ₂ C	0	CH ₂	1	=O	---	H	H	C	CH ₃	---
710	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	=O	---	H	H	A	CH ₃	---
711	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	=O	---	H	H	B	CH ₃	---
712	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	=O	---	H	H	C	CH ₃	---
713	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	H	H	H	H	A	CH ₃	---
714	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	H	H	H	H	B	CH ₃	---
715	N	[C ₂ H ₄ OC(=O)] ₂ C	1	CH ₂	1	H	H	H	H	C	CH ₃	---
716	N	(CH ₃ OCH ₂) ₂ C	1	CH ₂	1	H	H	H	H	A	CH ₃	---
717	N	(CH ₃ OCH ₂) ₂ C	1	CH ₂	1	H	H	H	H	B	CH ₃	---
718	N	(CH ₃ OCH ₂) ₂ C	1	CH ₂	1	H	H	H	H	C	CH ₃	---

Key: (a) R⁰ group codes: A =

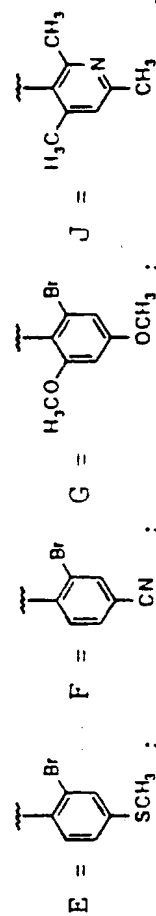
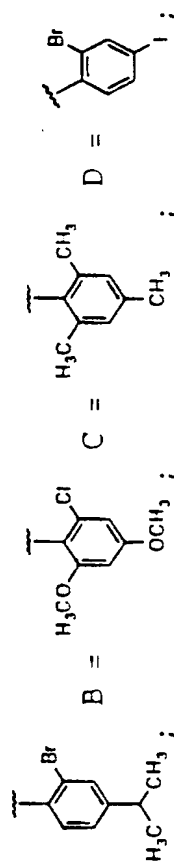
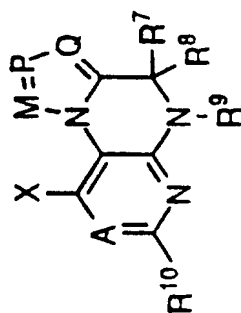


Table IV.



Ex. No.	A	X	M	P	Q	R ⁷	R ⁸	R ^{9 a}	R ¹⁰	m.p.
719	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	A	CH ₃	b
720	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	B	CH ₃	---
721	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	C	CH ₃	---
722	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	D	CH ₃	---
723	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	E	CH ₃	---
724	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	F	CH ₃	---
725	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	G	CH ₃	---
726	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	J	CH ₃	---
727	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	A	CH ₃	---
728	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	B	CH ₃	---
729	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	C	CH ₃	---
730	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	D	CH ₃	---
731	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	E	CH ₃	---
732	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	F	CH ₃	---
733	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	G	CH ₃	---
734	N	(C ₂ H ₅) ₂ N	N	N	N	H	H	J	CH ₃	---
735	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	A	CH ₃	---
736	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	B	CH ₃	---

737	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	C	CH ₃	---
738	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	D	CH ₃	---
739	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	E	CH ₃	---
740	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	F	CH ₃	---
741	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	G	CH ₃	---
742	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	J	CH ₃	---
743	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	A	CH ₃	---
744	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	B	CH ₃	---
745	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	C	CH ₃	---
746	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	D	CH ₃	---
747	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	E	CH ₃	---
748	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	F	CH ₃	---
749	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	G	CH ₃	---
750	N	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	J	CH ₃	---
751	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	A	CH ₃	---
752	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	B	CH ₃	---
753	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	C	CH ₃	---
754	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	D	CH ₃	---
755	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	E	CH ₃	---
756	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	F	CH ₃	---
757	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	G	CH ₃	---
758	N	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	J	CH ₃	---
759	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	A	CH ₃	---
760	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	B	CH ₃	---
761	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	C	CH ₃	---
762	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	D	CH ₃	---
763	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	E	CH ₃	---

764	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	F	CH ₃	---
765	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	G	CH ₃	---
766	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	J	CH ₃	---
767	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	A	CH ₃	---
768	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	B	CH ₃	---
769	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	C	CH ₃	---
770	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	D	CH ₃	---
771	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	E	CH ₃	---
772	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	F	CH ₃	---
773	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	G	CH ₃	---
774	N	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	J	CH ₃	---
775	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	A	CH ₃	---
776	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	B	CH ₃	---
777	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	C	CH ₃	---
778	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	D	CH ₃	---
779	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	E	CH ₃	---
780	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	F	CH ₃	---
781	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	G	CH ₃	---
782	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	J	CH ₃	---
783	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	A	CH ₃	---
784	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	B	CH ₃	---
785	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	C	CH ₃	---
786	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	D	CH ₃	---
787	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	E	CH ₃	---
788	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	F	CH ₃	---
789	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	G	CH ₃	---
790	N	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	J	CH ₃	---

791	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	A	CH ₃	---
792	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	B	CH ₃	---
793	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	C	CH ₃	---
794	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	D	CH ₃	---
795	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	E	CH ₃	---
796	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	F	CH ₃	---
797	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	G	CH ₃	---
798	N	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	J	CH ₃	---
799	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
800	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
801	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
802	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
803	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
804	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
805	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
806	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
807	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
808	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
809	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
810	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
811	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
812	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
813	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
814	N	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
815	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
816	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
817	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---

818	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
819	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
820	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
821	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
822	N	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
823	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
824	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
825	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
826	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
827	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
828	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
829	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
830	N	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
831	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
832	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
833	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
834	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
835	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
836	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
837	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
838	N	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
839	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	A	CH ₃	---
840	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	B	CH ₃	---
841	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	C	CH ₃	---
842	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	D	CH ₃	---
843	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	E	CH ₃	---
844	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	F	CH ₃	---

845	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	G	CH ₃	---
846	CH	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	J	CH ₃	---
847	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	A	CH ₃	---
848	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	B	CH ₃	---
849	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	C	CH ₃	---
850	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	D	CH ₃	---
851	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	E	CH ₃	---
852	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	F	CH ₃	---
853	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	G	CH ₃	---
854	CH	(C ₂ H ₅) ₂ N	N	N	N	H	H	J	CH ₃	---
855	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	A	CH ₃	---
856	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	B	CH ₃	---
857	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	C	CH ₃	---
858	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	D	CH ₃	---
859	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	E	CH ₃	---
860	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	F	CH ₃	---
861	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	G	CH ₃	---
862	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	N	N	N	H	H	J	CH ₃	---
863	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	A	CH ₃	---
864	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	B	CH ₃	---
865	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	C	CH ₃	---
866	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	D	CH ₃	---
867	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	E	CH ₃	---
868	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	F	CH ₃	---
869	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	G	CH ₃	---
870	CH	[CH ₃ O(CH ₂) ₂] ₂ N	N	N	N	H	H	J	CH ₃	---
871	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	A	CH ₃	---

872	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	B	CH ₃	---
873	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	C	CH ₃	---
874	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	D	CH ₃	---
875	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	E	CH ₃	---
876	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	F	CH ₃	---
877	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	G	CH ₃	---
878	CH	(C ₂ H ₅) ₂ CHNH	N	N	N	H	H	J	CH ₃	---
879	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	A	CH ₃	---
880	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	B	CH ₃	---
881	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	C	CH ₃	---
882	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	D	CH ₃	---
883	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	E	CH ₃	---
884	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	F	CH ₃	---
885	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	G	CH ₃	---
886	CH	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	J	CH ₃	---
887	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	A	CH ₃	---
888	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	B	CH ₃	---
889	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	C	CH ₃	---
890	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	D	CH ₃	---
891	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	E	CH ₃	---
892	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	F	CH ₃	---
893	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	G	CH ₃	---
894	CH	(C ₂ H ₅) ₂ N	CH	CH	N	H	H	J	CH ₃	---
895	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	A	CH ₃	---
896	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	B	CH ₃	---
897	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	C	CH ₃	---
898	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	D	CH ₃	---

899	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	E	CH ₃	---
900	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	F	CH ₃	---
901	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	G	CH ₃	---
902	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CH	CH	N	H	H	J	CH ₃	---
903	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	A	CH ₃	---
904	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	B	CH ₃	---
905	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	C	CH ₃	---
906	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	D	CH ₃	---
907	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	E	CH ₃	---
908	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	F	CH ₃	---
909	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	G	CH ₃	---
910	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CH	CH	N	H	H	J	CH ₃	---
911	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	A	CH ₃	---
912	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	B	CH ₃	---
913	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	C	CH ₃	---
914	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	D	CH ₃	---
915	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	E	CH ₃	---
916	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	F	CH ₃	---
917	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	G	CH ₃	---
918	CH	(C ₂ H ₅) ₂ CHNH	CH	CH	N	H	H	J	CH ₃	---
919	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
920	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
921	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
922	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
923	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
924	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
925	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---

926	CH	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
927	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
928	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
929	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
930	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
931	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
932	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
933	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
934	CH	(C ₂ H ₅) ₂ N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
935	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
936	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
937	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
938	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
939	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
940	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
941	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
942	CH	CH ₃ OCH ₂ CH(C ₂ H ₅)NH	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
943	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
944	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---
945	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
946	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
947	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
948	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
949	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
950	CH	[CH ₃ O(CH ₂) ₂] ₂ N	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
951	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	A	CH ₃	---
952	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	B	CH ₃	---

953	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	C	CH ₃	---
954	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	D	CH ₃	---
955	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	E	CH ₃	---
956	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	F	CH ₃	---
957	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	G	CH ₃	---
958	CH	(C ₂ H ₅) ₂ CHNH	CCH ₃	CCH ₃	N	H	H	J	CH ₃	---
959	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	CH ₃	CH ₃	A	CH ₃	---
960	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	A	H	---
961	N	C ₄ H ₉ (C ₂ H ₅)N	N	N	N	H	H	A	CN	---
962	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	CH ₃	CH ₃	A	CH ₃	---
963	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	A	H	---
964	N	C ₄ H ₉ (C ₂ H ₅)N	CH	CH	N	H	H	A	CN	---
965	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	CH ₃	CH ₃	A	CH ₃	---
966	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	A	H	---
967	N	C ₄ H ₉ (C ₂ H ₅)N	CCH ₃	CCH ₃	N	H	H	A	CN	---

UtilityCRF-R1 Receptor Binding Assay for the Evaluation of
Biological Activity

5

The following is a description of the isolation of cell membranes containing cloned human CRF-R1 receptors for use in the standard binding assay as well as a description of the assay itself.

10 Messenger RNA was isolated from human hippocampus. The mRNA was reverse transcribed using oligo (dt) 12-18 and the coding region was amplified by PCR from start to stop codons. The resulting PCR fragment was cloned into the EcoRV site of pGEMV, from whence the insert was
15 reclaimed using XhoI + XbaI and cloned into the XhoI + XbaI sites of vector pm3ar (which contains a CMV promoter, the SV40 't' splice and early poly A signals, an Epstein-Barr viral origin of replication, and a hygromycin selectable marker). The resulting expression
20 vector, called phchCRFR was transfected in 293EBNA cells and cells retaining the episome were selected in the presence of 400 μ M hygromycin. Cells surviving 4 weeks of selection in hygromycin were pooled, adapted to growth in suspension and used to generate membranes for
25 the binding assay described below. Individual aliquots containing approximately 1×10^8 of the suspended cells were then centrifuged to form a pellet and frozen.

For the binding assay a frozen pellet described above containing 293EBNA cells transfected with hCRFR1
30 receptors is homogenized in 10 ml of ice cold tissue buffer (50 mM HEPES buffer pH 7.0, containing 10 mM $MgCl_2$, 2 mM EGTA, 1 μ g/ml aprotinin, 1 μ g/ml leupeptin and 1 μ g/ml pepstatin). The homogenate is centrifuged at 40,000 x g for 12 min and the resulting pellet
35 rehomogenized in 10 ml of tissue buffer. After another centrifugation at 40,000 x g for 12 min, the pellet is

resuspended to a protein concentration of 360 µg/ml to be used in the assay.

Binding assays are performed in 96 well plates; each well having a 300 µl capacity. To each well is added 50 µl of test drug dilutions (final concentration of drugs range from 10^{-10} - 10^{-5} M), 100 µl of ^{125}I -ovine-CRF (^{125}I -o-CRF) (final concentration 150 pM) and 150 µl of the cell homogenate described above. Plates are then allowed to incubate at room temperature for 2 hours before filtering the incubate over GF/F filters (presoaked with 0.3% polyethyleneimine) using an appropriate cell harvester. Filters are rinsed 2 times with ice cold assay buffer before removing individual filters and assessing them for radioactivity on a gamma counter.

Curves of the inhibition of ^{125}I -o-CRF binding to cell membranes at various dilutions of test drug are analyzed by the iterative curve fitting program LIGAND (P.J. Munson and D. Rodbard, *Anal. Biochem.* 107:220 (1980)), which provides K_i values for inhibition which are then used to assess biological activity.

A compound is considered to be active if it has a K_i value of less than about 10000 nM for the inhibition of CRF.

Inhibition of CRF-Stimulated Adenylate Cyclase Activity

Inhibition of CRF-stimulated adenylate cyclase activity was performed as described by G. Battaglia et al. *Synapse* 1:572 (1987). Briefly, assays were carried out at 37° C for 10 min in 200 µl of buffer containing 100 mM Tris-HCl (pH 7.4 at 37° C), 10 mM MgCl_2 , 0.4 mM EGTA, 0.1% BSA, 1 mM isobutylmethylxanthine (IBMX), 250 units/ml phosphocreatine kinase, 5 mM creatine phosphate, 100 mM guanosine 5'-triphosphate, 100 nM oCRF, antagonist peptides (concentration range 10^{-9} to 10^{-6} M) and 0.8 mg original wet weight tissue (approximately 40-60 µg protein). Reactions were

initiated by the addition of 1 mM ATP/³²P]ATP (approximately 2-4 mCi/tube) and terminated by the addition of 100 ml of 50 mM Tris-HCL, 45 mM ATP and 2% sodium dodecyl sulfate. In order to monitor the recovery of cAMP, 1 µl of [³H]cAMP (approximately 40,000 dpm) was added to each tube prior to separation. The separation of [³²P]cAMP from [³²P]ATP was performed by sequential elution over Dowex and alumina columns. Recovery was consistently greater than 80%.

Some compounds of this invention were tested in this assay and found to be active.

In vivo Biological Assay

The *in vivo* activity of the compounds of the present invention can be assessed using any one of the biological assays available and accepted within the art. Illustrative of these tests include the Acoustic Startle Assay, the Stair Climbing Test, and the Chronic Administration Assay. These and other models useful for the testing of compounds of the present invention have been outlined in C.W. Berridge and A.J. Dunn *Brain Research Reviews* 15:71 (1990)

Compounds may be tested in any species of rodent or small mammal. Disclosure of the assays herein is not intended to limit the enablement of the invention.

Compounds of this invention have utility in the treatment of imbalances associated with abnormal levels of corticotropin releasing factor in patients suffering from depression, affective disorders, and/or anxiety.

Compounds of this invention can be administered to treat these abnormalities by means that produce contact of the active agent with the agent's site of action in the body of a mammal. The compounds can be administered by any conventional means available for use in conjunction with pharmaceuticals either as individual therapeutic agent or in combination of therapeutic agents. They can be administered alone, but will

generally be administered with a pharmaceutical carrier selected on the basis of the chosen route of administration and standard pharmaceutical practice.

The dosage administered will vary depending on the use and known factors such as pharmacodynamic character of the particular agent, and its mode and route of administration; the recipient's age, weight, and health; nature and extent of symptoms; kind of concurrent treatment; frequency of treatment; and desired effect.

For use in the treatment of said diseases or conditions, the compounds of this invention can be orally administered daily at a dosage of the active ingredient of 0.002 to 200 mg/kg of body weight. Ordinarily, a dose of 0.01 to 10 mg/kg in divided doses one to four times a day, or in sustained release formulation will be effective in obtaining the desired pharmacological effect.

Dosage forms (compositions) suitable for administration contain from about 1 mg to about 100 mg of active ingredient per unit. In these pharmaceutical compositions, the active ingredient will ordinarily be present in an amount of about 0.5 to 95% by weight based on the total weight of the composition.

The active ingredient can be administered orally is solid dosage forms, such as capsules, tablets and powders; or in liquid forms such as elixirs, syrups, and/or suspensions. The compounds of this invention can also be administered parenterally in sterile liquid dose formulations.

Gelatin capsules can be used to contain the active ingredient and a suitable carrier such as but not limited to lactose, starch, magnesium stearate, steric acid, or cellulose derivatives. Similar diluents can be used to make compressed tablets. Both tablets and capsules can be manufactured as sustained release products to provide for continuous release of medication over a period of time. Compressed tablets can be sugar-

coated or film-coated to mask any unpleasant taste, or used to protect the active ingredients from the atmosphere, or to allow selective disintegration of the tablet in the gastrointestinal tract.

- 5 Liquid dose forms for oral administration can contain coloring or flavoring agents to increase patient acceptance.

- 10 In general, water, pharmaceutically acceptable oils, saline, aqueous dextrose (glucose), and related sugar solutions and glycols, such as propylene glycol or polyethylene glycol, are suitable carriers for parenteral solutions. Solutions for parenteral administration preferably contain a water soluble salt of the active ingredient, suitable stabilizing agents, and if necessary, buffer substances. Antioxidizing agents, such as sodium bisulfite, sodium sulfite, or ascorbic acid, either alone or in combination, are suitable stabilizing agents. Also used are citric acid and its salts, and EDTA. In addition, parenteral solutions can contain preservatives such as benzalkonium chloride, methyl- or propyl-paraben, and chlorobutanol.

- 20 Suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences", A. Osol, a standard reference in the field.

- 25 Useful pharmaceutical dosage-forms for administration of the compounds of this invention can be illustrated as follows:

Capsules

- 30 A large number of units capsules are prepared by filling standard two-piece hard gelatin capsules each with 100 mg of powdered active ingredient, 150 mg lactose, 50 mg cellulose, and 6 mg magnesium stearate.

35 Soft Gelatin Capsules

- A mixture of active ingredient in a digestible oil such as soybean, cottonseed oil, or olive oil is

prepared and injected by means of a positive displacement was pumped into gelatin to form soft gelatin capsules containing 100 mg of the active ingredient. The capsules were washed and dried.

5

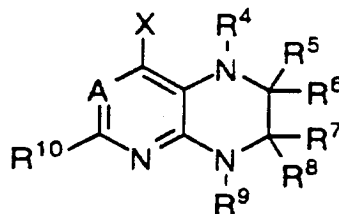
Tablets

A large number of tablets are prepared by conventional procedures so that the dosage unit was 100 mg active ingredient, 0.2 mg of colloidal silicon dioxide, 5 mg of magnesium stearate, 275 mg of microcrystalline cellulose, 11 mg of starch, and 98.8 mg lactose. Appropriate coatings may be applied to increase palatability or delayed adsorption.

15 The compounds of this invention may also be used as reagents or standards in the biochemical study of neurological function, dysfunction, and disease.

Claims:

1. A method of treating an affective disorder,
 5 anxiety, depression, irritable bowel syndrome, post-traumatic stress disorder, supranuclear palsy, immune suppression, Alzheimer's disease, gastrointestinal disease, anorexia nervosa or other feeding disorder,
 10 drug or alcohol withdrawal symptoms, drug addiction, or inflammatory disorder in a mammal comprising administering to the mammal a therapeutically effective amount of a CRF antagonist compound of formula I:



(I)

or a pharmaceutically acceptable salt or prodrug thereof, wherein:

- A is N or C-R¹¹;
 X is H, OR¹, S(O)_nR¹, NR¹R², CR¹R²R³, phenyl (optionally
 20 substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl) or heteroaryl (optionally
 25 substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
 30 n is 0, 1 or 2;
 R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, C₂-C₁₂ alkenyl, C₂-C₁₂ alkynyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂

dialkylaminoalkyl, C₂-C₁₃ cyanoalkyl, C₂-C₅ carboalkoxy-(C₁-C₁₂ alkyl), phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl), or heteroaryl (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);

R² and R³ are independently chosen from H, C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, C₂-C₁₂ alkenyl, C₂-C₁₂ alkynyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, C₂-C₁₃ cyanoalkyl, C₁-C₄ carboalkoxy, C₂-C₁₂ carboalkoxyalkyl, C(=O)CH₃, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl), or heteroaryl (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);

R⁴ is H, C₁-C₁₂ alkyl, allyl, propargyl or benzyl (optionally substituted with 1-4 groups independently chosen from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);

R¹ and R⁴ may also optionally be taken together, along with the other four interconnected atoms, to form a ring of 5-9 total atoms, the structural sequence

between the X group and the ring nitrogen atom consisting of the group $(CH_2)_pW(CH_2)_q$;

p and q are independently 0, 1 or 2;

W is CH_2 , $C(CH_3)_2$, $C(=O)$, O, S or NCH_3 ;

5 R^5 , R^6 , R^7 and R^8 are independently chosen from H,

C_1 - C_4 alkyl, allyl, propargyl, phenyl (optionally substituted with 1-4 groups independently chosen

from halogen, C_1 - C_4 haloalkyl, nitro, C_1 - C_4 alkyl,

C_2 - C_5 carboalkoxy, cyano, OH, C_1 - C_4 alkoxy, SH, C_1 - C_4

10 alkylthio, NH_2 , C_1 - C_4 alkylamino, C_2 - C_8

dialkylamino, or phenyl) or benzyl (optionally

substituted with 1-4 groups independently chosen

from halogen, C_1 - C_4 haloalkyl, nitro, C_1 - C_4 alkyl,

C_2 - C_5 carboalkoxy, cyano, OH, C_1 - C_4 alkoxy, SH, C_1 - C_4

15 alkylthio, NH_2 , C_1 - C_4 alkylamino, C_2 - C_8

dialkylamino, or phenyl);

R^4 , R^5 and R^6 may also be taken together, along with the two interconnecting atoms, to constitute either

an imidazole or tetrazole ring, the imidazole ring

20 being optionally substituted with 1-2 groups chosen independently from C_1 - C_4 alkyl or phenyl;

R^5 and R^6 may also be taken together to be O, S or NR^{12} ;

R^9 is phenyl (optionally substituted with 1-4 groups

25 chosen from halogen, C_1 - C_4 haloalkyl, C_1 - C_4 alkyl,

C_2 - C_6 alkenyl, C_1 - C_4 alkoxy, C_1 - C_4 alkylthio, C_1 - C_4

alkylsulfonyl, C_2 - C_6 dialkylamino, nitro, C_2 - C_5

carboalkoxy or cyano), pyridyl (optionally

substituted with 1-4 groups chosen from halogen,

30 C_1 - C_4 haloalkyl, C_1 - C_4 alkyl, C_2 - C_6 alkenyl, C_1 - C_4

alkoxy, C_1 - C_4 alkylthio, C_1 - C_4 alkylsulfonyl, C_2 - C_6

dialkylamino, nitro, C_2 - C_5 carboalkoxy or cyano),

or pyrimidyl (optionally substituted with 1-4

35 groups chosen from halogen, C_1 - C_4 haloalkyl, C_1 - C_4

alkyl, C_2 - C_6 alkenyl, C_1 - C_4 alkoxy, C_1 - C_4

alkylthio, C_1 - C_4 alkylsulfonyl, C_2 - C_6 dialkylamino,

nitro, C_2 - C_5 carboalkoxy or cyano);

R¹⁰ is H, C₁-C₄ alkyl or cyano;

R¹¹ is H, C₁-C₄ alkyl or halogen;

R¹² is H, C₁-C₄ alkyl or phenyl;

aryl is phenyl, biphenyl or naphthyl; and

5 heteroaryl is pyridyl, pyrimidinyl, triazinyl, furanyl, quinolinyl, isoquinolinyl, thienyl, imidazolyl, thiazolyl, indolyl, pyrrolyl, oxazolyl, benzofuranyl, benzothienyl, benzthiazolyl, isoxazolyl or pyrazolyl.

10

2. Method of claim 1 wherein, in the compound of formula I,

X is OR¹, NR¹R², CR¹R²R³, or phenyl (optionally substituted at the 2-position with CF₃, nitro, halogen or cyano);

15

R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, or phenyl (optionally substituted with 1-4 groups independently chosen from halogen, haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₆ dialkylamino, or phenyl);

20

R⁴ is H or C₁-C₄ alkyl;

25

R⁵ and R⁶ are either H or C₁-C₄ alkyl;

R⁴, R⁵ and R⁶ may also be taken together, along with the two interconnecting atoms, to constitute a tetrazole ring;

30

R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), 3-pyridyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano),

35

or 5-pyrimidyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₆ carboalkoxy or cyano);

R¹⁰ is CH₃;

and R¹¹ is H.

3. Method of claim 2 wherein, in the compound of formula (I):

A is N;

X is NR¹R² or CR¹R²R³;

R¹ is C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;

R² and R³ are independently H, C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;

R⁴ is H;

R⁵ and R⁶ are H;

R⁷ and R⁸ are independently H or CH₃; and

R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₆ carboalkoxy or cyano).

4. Method of claim 1 wherein the compound of formula I is selected from:

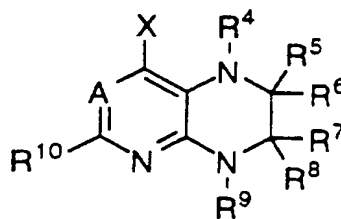
8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;

8-(2-chloro-4,6-dimethoxyphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;

4-(ethylbutylamino)-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine; and

4-(1-methoxy-2-butyl)amino-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine.

5. A CRF antagonist compound of formula I:



(I)

or a pharmaceutically acceptable salt or prodrug thereof,
wherein:

- 5 A is N or C-R¹¹;
 X is H, OR¹, S(O)_nR¹, NR¹R², CR¹R²R³, phenyl (optionally
 substituted with 1-4 groups independently chosen
 from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl,
 C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄
 10 alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈
 dialkylamino, or phenyl) or heteroaryl (optionally
 substituted at one to all valence-allowed positions
 with groups independently chosen from halogen, C₁-
 C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy,
 15 cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂,
 C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);
 n is 0, 1 or 2;
 R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂
 cycloalkyl, C₄-C₁₂ cycloalkylalkyl, C₂-C₁₂ alkenyl,
 20 C₂-C₁₂ alkynyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂
 dialkylaminoalkyl, C₂-C₁₃ cyanoalkyl, C₂-C₅
 carboalkoxy-(C₁-C₁₂ alkyl), phenyl (optionally
 substituted with 1-4 groups independently chosen
 from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl,
 25 C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄
 alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₈
 dialkylamino, or phenyl); or heteroaryl (optionally
 substituted at one to all valence-allowed positions
 with groups independently chosen from halogen, C₁-
 30 C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy,
 cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂,
 C₁-C₄ alkylamino, C₂-C₈ dialkylamino, or phenyl);

R^2 and R^3 are independently chosen from H, C_1-C_{12} alkyl, C_2-C_{12} alkoxyalkyl, C_3-C_{12} cycloalkyl, C_4-C_{12} cycloalkylalkyl, C_2-C_{12} alkenyl, C_2-C_{12} alkynyl, aryl- (C_1-C_{12} alkyl), C_3-C_{12} dialkylaminoalkyl, C_2-C_{13} cyanoalkyl, C_1-C_4 carboalkoxy, C_2-C_{12} carboalkoxyalkyl, $C(=O)CH_3$, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C_1-C_4 haloalkyl, nitro, C_1-C_4 alkyl, C_2-C_5 carboalkoxy, cyano, OH, C_1-C_4 alkoxy, SH, C_1-C_4 alkylthio, NH_2 , C_1-C_4 alkylamino, C_2-C_8 dialkylamino, or phenyl), or heteroaryl' (optionally substituted at one to all valence-allowed positions with groups independently chosen from halogen, C_1-C_4 haloalkyl, nitro, C_1-C_4 alkyl, C_2-C_5 carboalkoxy, cyano, OH, C_1-C_4 alkoxy, SH, C_1-C_4 alkylthio, NH_2 , C_1-C_4 alkylamino, C_2-C_8 dialkylamino, or phenyl);

R^4 is H, C_1-C_{12} alkyl, allyl, propargyl or benzyl (optionally substituted with 1-4 groups independently chosen from halogen, C_1-C_4 haloalkyl, nitro, C_1-C_4 alkyl, C_2-C_5 carboalkoxy, cyano, OH, C_1-C_4 alkoxy, SH, C_1-C_4 alkylthio, NH_2 , C_1-C_4 alkylamino, C_2-C_8 dialkylamino, or phenyl);

R^1 and R^4 may also optionally be taken together, along with the other four interconnected atoms, to form a ring of 5-9 total atoms, the structural sequence between the X group and the ring nitrogen atom consisting of the group $(CH_2)_pW(CH_2)_q$;

p and q are independently 0, 1 or 2;
W is CH_2 , $C(CH_3)_2$, $C(=O)$, O, S or NCH_3 ;

R^5 , R^6 , R^7 and R^8 are independently chosen from H, C_1-C_4 alkyl, allyl, propargyl, phenyl (optionally substituted with 1-4 groups independently chosen from halogen, C_1-C_4 haloalkyl, nitro, C_1-C_4 alkyl, C_2-C_5 carboalkoxy, cyano, OH, C_1-C_4 alkoxy, SH, C_1-C_4 alkylthio, NH_2 , C_1-C_4 alkylamino, C_2-C_8 dialkylamino, or phenyl) or benzyl (optionally substituted with 1-4 groups independently chosen

from halogen, C₁-C₄ haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₅ dialkylamino, or phenyl);

5 R⁴, R⁵ and R⁶ may also be taken together, along with the two interconnecting atoms, to constitute either an imidazole or tetrazole ring, the imidazole ring being optionally substituted with 1-2 groups chosen independently from C₁-C₄ alkyl or phenyl;

10 R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), pyridyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), or pyrimidyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano);

R¹⁰ is H, C₁-C₄ alkyl or cyano;

25 R¹¹ is H, C₁-C₄ alkyl or halogen;

R¹² is H, C₁-C₄ alkyl or phenyl;

aryl is phenyl, biphenyl or naphthyl; and

heteroaryl is pyridyl, pyrimidinyl, triazinyl,

furanyl, quinolinyl, isoquinolinyl, thienyl,

30 imidazolyl, thiazolyl, indolyl, pyrrolyl, oxazolyl, benzofuranyl, benzothienyl, benzthiazolyl, isoxazolyl or pyrazolyl.

6. A compound of claim 5 wherein:

35

X is OR¹, NR¹R², CR¹R²R³, or phenyl (optionally

- substituted at the 2-position with CF₃, nitro, halogen or cyano);
- 5 R¹ is C₁-C₁₂ alkyl, C₂-C₁₂ alkoxyalkyl, C₃-C₁₂ cycloalkyl, C₄-C₁₂ cycloalkylalkyl, aryl-(C₁-C₁₂ alkyl), C₃-C₁₂ dialkylaminoalkyl, or phenyl (optionally substituted with 1-4 groups independently chosen from halogen, haloalkyl, nitro, C₁-C₄ alkyl, C₂-C₅ carboalkoxy, cyano, OH, C₁-C₄ alkoxy, SH, C₁-C₄ alkylthio, NH₂, C₁-C₄ alkylamino, C₂-C₆ dialkylamino, or phenyl);
- 10 R⁴ is H or C₁-C₄ alkyl;
- R⁵ and R⁶ are either H or C₁-C₄ alkyl;
- R⁴, R⁵ and R⁶ may also be taken together, along with the two interconnecting atoms, to constitute a
- 15 tetrazole ring;
- R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano), 3-pyridyl (optionally
- 20 substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano),
- 25 or 5-pyrimidyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano);
- 30 R¹⁰ is CH₃;
- and R¹¹ is H.

7. A compound of claim 6 wherein:

- 35 A is N;
- X is NR¹R² or CR¹R²R³;
- R¹ is C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;

- R² and R³ are independently H, C₁-C₆ alkyl or C₂-C₈ alkoxyalkyl;
R⁴ is H;
R⁵ and R⁶ are H;
5 R⁷ and R⁸ are independently H or CH₃; and
R⁹ is phenyl (optionally substituted with 1-4 groups chosen from halogen, C₁-C₄ haloalkyl, C₁-C₄ alkyl, C₂-C₆ alkenyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfonyl, C₂-C₆ dialkylamino, nitro, C₂-C₅ carboalkoxy or cyano).
10

8. A compound of claim 7 selected from:

- 8-(2-bromo-4-isopropylphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;
15 8-(2-chloro-4,6-dimethoxyphenyl)-4-(ethylbutylamino)-2-methyl-5,6,7,8-tetrahydropteridine;
4-(ethylbutylamino)-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine; and
20 4-(1-methoxy-2-butyl)amino-2-methyl-8-(2,4,6-trimethylphenyl)-5,6,7,8-tetrahydropteridine.

9. A composition comprising a therapeutically effective amount of compound of claim 5 and a
25 pharmaceutically suitable carrier.

10. A composition comprising a therapeutically effective amount of compound of claim 6 and a
30 pharmaceutically suitable carrier.

11. A composition comprising a therapeutically effective amount of compound of claim 7 and a
pharmaceutically suitable carrier.

35 12. A composition comprising a therapeutically effective amount of compound of claim 8 and a pharmaceutically suitable carrier.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/08448

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 31/495, 31/505; C07D 487/06, 487/16

US CL : 514/249; 544/251, 256, 350

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/249; 544/258, 260, 251, 256

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

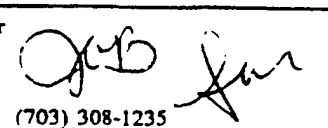
CAS ONLINE STRUCTURE SEARCH

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X - Y	US 5,198,547 A (BAILEY et al.) 30 March 1993, see column 7 structure IX and column 2 lines 64 to column 3 line 14.	5, 6, 9, 10 -----1, 2
A	US 4,665,182 A (NICHOL et al.) 12 May 1987, see claim 1	1-12
A	US 4,701,455 A (NICHOL et al.) 20 October 1987, see claim 1.	1-12
A	US 4,091,219 A (DENZEL et al.) 23 May 1978, see Table, columns 5 to 8.	1-12
X	US 4,510,141 A (HECKENDORN) 09 April 1985, see column 20 line 34 to column 25 line 8 and Abstract Penultimate line.	1, 2, 5, 6, 9, 10

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A document defining the general state of the art which is not considered to be of particular relevance	*X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E earlier document published on or after the international filing date	*Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z document member of the same patent family
*O document referring to an oral disclosure, use, exhibition or other means	
*P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 21 AUGUST 1997	Date of mailing of the international search report 09 SEP 1997
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer D. G. DAUS  Telephone No. (703) 308-1235

Form PCT/ISA/210 (second sheet)(July 1992)*

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/08448

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JP 08-092250 (A. SAHI BREWERIES) 09 April 1996, see page 2, column 1 structure (3) and column 2, structure (1).	1-12

Form PCT/ISA/210 (continuation of second sheet)(July 1992)*

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/08448

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

- I. Claims 1-2, (in part) 3, 4 to various disease treatments using III (A is N)
- II. Claims 1-2 (in part) to various disease treatments using IV (a is CH)
- III. Claims 5, 6, 9, 10 (in part) and 7, 8, 11, 12 to tetrahydropteridine (A is N)
- IV. Claims 5, 6, 9, 10 (in part) to pyridopiperazines (A is CH).

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.